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Title: COMPOUNDS AND METHODS TO INHIBIT OR AUGMENT AN INFLAMMATORY RESPONSE

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CONTINUATION-IN-PART (CIP) of prior Patent Application No. 08/927,939 (under 37 CFR § 1.53(b)) comprising:

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 Formal Drawing(s) (24 sheets).
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**COMPOUNDS AND METHODS TO INHIBIT
OR AUGMENT AN INFLAMMATORY RESPONSE**

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Cross-Reference to Related Applications

This application is a continuation-in-part application of U.S. application Serial No. 08/927,939, filed September 11, 1997, the disclosure of which is incorporated by reference herein.

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Background of the Invention

Macrophage/monocyte recruitment plays a role in the morbidity and mortality of a broad spectrum of diseases, including autoimmune diseases, granulomatous diseases, allergic diseases, infectious diseases, osteoporosis and coronary artery disease. For example, in atherosclerosis early during lipid lesion formation, circulating monocytes adhere to the activated endothelium overlying the incipient plaque. Under appropriate conditions, the monocytes then migrate into the developing intima. In the intima, macrophage accumulate lipoprotein and excrete an excess of proteases relative to protease inhibitors. If the lipoproteins are oxidized, they are toxic to macrophage, which results in macrophage death and an increase in an unstable, necrotic, extracellular lipid pool. An excess of proteases results in loss of extracellular matrix and destabilization of the fibrous plaque. Plaque instability is the acute cause of myocardial infarction.

Many molecules have been identified that are necessary for the recruitment of monocytes and other inflammatory cell types. These molecules represent targets for the inhibition of monocyte recruitment. One class of such molecules is adhesion molecules, e.g., receptors, for monocytes. Another class of molecules includes inflammatory mediators, such as TNF- α and related molecules, the interleukins, e.g., IL-1 β , and chemokines, e.g., monocyte chemoattractant protein-1 (MCP-1). As a result, agents which modulate the activity of chemokines are likely to be useful to prevent and treat a wide range of diseases. For example, Rollins et al. (U.S. Patent No. 5,459,128) generally disclose analogs of MCP-1 that inhibit the

monocyte chemoattractant activity of endogenous MCP-1. Analogs that are effective to inhibit endogenous MCP-1 are disclosed as analogs which are modified at 28-tyrosine, 24-arginine, 3-aspartate and/or in amino acids between residues 2-8 of MCP-1. In particular, Rollins et al. state that “[s]uccessful inhibition of the 5 activity is found where MCP-1 is modified in one or more of the following ways: a) the 28-tyrosine is substituted by aspartate, b) the 24-arginine is substituted by phenylalanine, c) the 3-aspartate is substituted by alanine, and/or d) the 2-8 amino acid sequence is deleted” (col. 1, lines 49-54). The deletion of amino acids 2-8 of MCP-1 (“MCP-1(Δ 2-8)”) results in a polypeptide that is inactive, i.e., MCP-1(Δ 2-8) 10 is not a chemoattractant (col. 5, lines 22-23). The only effective MCP-1 inhibitor disclosed in Rollins et al. is MCP-1(Δ 2-8).

Recent studies suggest that MCP-1(Δ 2-8) exhibits a dominant negative effect, i.e., it forms heterodimers with wild-type MCP-1 that cannot elicit a biological effect (Zhang et al., J. Biol. Chem., 269, 15918 (1994); Zhang et al., Mol. 15 Cell. Biol., 15, 4851 (1995)). Thus, MCP-1(Δ 2-8) does not exhibit properties of a classic receptor antagonist. Moreover, MCP-1(Δ 2-8) is unlikely to be widely useful for inhibition of MCP-1 activity *in vivo*, as MCP-1(Δ 2-8) is a large polypeptide with undesirable pharmacodynamic properties. Furthermore, it is unknown whether MCP-1(Δ 2-8) is active as a dominant-negative inhibitor of other chemokines 20 associated with inflammation.

Thus, there is a need to identify agents that inhibit or enhance chemokine-induced macrophage and/or monocyte recruitment and which have desirable pharmacodynamic properties. Moreover, there is a need to identify agents that inhibit or enhance chemokine-induced activities of other cell types, such as 25 lymphocytes. Further, there is a need to identify agents that are pan-selective chemokine inhibitors.

Summary of the Invention

The invention provides a therapeutic agent comprising an isolated and purified chemokine peptide, chemokine peptide variant, chemokine analog, or a derivative thereof. Preferably, the therapeutic agent of the invention inhibits the 5 activity of more than one chemokine, although the agent may not inhibit the activity of all chemokines to the same extent. Alternatively, a preferred therapeutic agent of the invention specifically inhibits the activity of one chemokine to a greater extent than other chemokines. Yet another preferred therapeutic agent of the invention mimics the activity of a chemokine, e.g., it acts as an agonist. Thus, therapeutic 10 agents that are chemokine antagonists and agonists are within the scope of the invention. A further preferred therapeutic agent of the invention is an agent that does not inhibit or mimic the activity of a chemokine but binds to or near the receptor for that chemokine, i.e., it is a neutral agent.

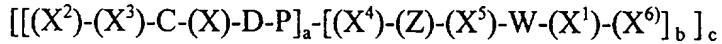
A preferred embodiment of the invention is an isolated and purified CC 15 chemokine peptide 3, e.g., a peptide derived from MCP-1 which corresponds to about residue 46 to about residue 67 of mature MCP-1 (“peptide 3[MCP-1”]), a variant, an analog, or a derivative thereof. It is contemplated that chemokine peptide 3, a variant, an analog or a derivative thereof is a chemokine receptor antagonist, although these therapeutic agents may exert their effect by a different mechanism, 20 e.g., by inhibiting the arachidonic acid pathway (e.g., inhibition of leukotriene, thromboxane, or prostaglandin synthesis or stability) or by elevating TGF-beta levels, or by more than one mechanism.

A preferred peptide 3 of the invention is a compound of formula (I):



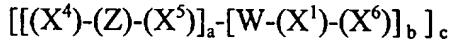
25 wherein X^2 is E, Q, D, N, L, P, I or M, wherein X^3 is I, V, M, A, P, norleucine or L, wherein X is A, L, V, M, P, norleucine or I, wherein X^4 is K, S, R, R, Q, N or T, wherein Z is Q, K, E, N, R, I, V, M, A, P, norleucine or L, wherein X^7 is D or P, wherein X^5 is K, E, R, S, Q, D, T, H or N, wherein X^1 is V, L, M, P, A, norleucine, or I, wherein X^6 is Q, N, K or R, wherein a is 0-6, wherein b is 0-6, and wherein c is

1-6, with the proviso that a and b cannot both be 0. The letters in formulas (I)-(III) that are not X, Y or Z represent peptidyl residues as shown in Figure 13. A more preferred peptide 3 of the invention is a compound of formula (I):



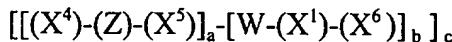
5 wherein X^2 is E, Q or M, wherein X^3 is I, V or L, wherein X is A, L or I, wherein X^4 is K, S or T, wherein Z is Q, K, E or L, wherein X^5 is K, E, R, S or T, wherein X^1 is V or I, wherein X^6 is Q or R, wherein a is 0-6, wherein b is 0-6, and wherein c is 1-6, with the proviso that a and b cannot both be 0.

10 Yet another preferred peptide 3 of the invention is a compound of formula (II):



wherein X^4 is K, S or T, wherein Z is Q, K, E or L, wherein X^5 is K, E, R, S or T, wherein X^1 is V or I, wherein X^6 is Q or R, wherein a is 0-6, wherein b is 0-6, and wherein c is 1-6, with the proviso that a and b cannot both be 0.

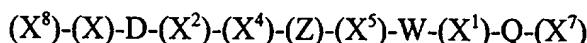
15 Another preferred peptide 3 of the invention is a compound of formula (III):



wherein X^4 is K, S, R, R, Q, N or T, wherein Z is Q, K, E, N, R, I, V, M, A, P, norleucine or L, wherein X^5 is K, E, R, S, Q, D, T, H or N, wherein X^1 is V, L, M, P, A, norleucine, or I, wherein X^6 is Q, N, K or R, wherein a is 0-6, wherein b is 0-6,

20 and wherein c is 1-6, with the proviso that a and b cannot both be 0.

A more preferred peptide 3 of the invention is a compound of formula (X):



wherein X is A, L, V or I, wherein X^2 is P, G or L, wherein X^4 is K, T, R or N, wherein Z is Q, K, A or L, wherein X^5 is K, E, R, Q or P, wherein X^1 is V, L, A, M,

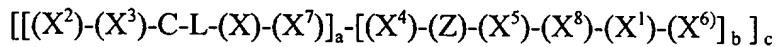
25 F or I, and wherein X^8 and X^7 are independently C or absent.

A preferred embodiment of the invention is an isolated and purified CC chemokine peptide 3, e.g., a peptide derived from MCP-1 which corresponds to SEQ ID NO:1 (“peptide 3(1-12)[MCP-1]”) or SEQ ID NO:7 (“peptide 3(3-12)[MCP-1]”), a fragment, a variant, an analog, or a derivative thereof. As

described hereinbelow, peptide 3(1-12)[MCP-1](SEQ ID NO:1) and peptide 3(3-12)[MCP-1] (SEQ ID NO:7) are pan-chemokine inhibitors, bioavailable, and have desirable pharmacokinetics. Another preferred CC chemokine peptide 3 of the invention is peptide 3[MIP1 α], and more preferably peptide 3(1-12)[MIP1 α] which 5 has an amino acid sequence corresponding to SEQ ID NO:42, a variant, an analog, a fragment or a derivative thereof.

Further preferred embodiments of the invention are a CC chemokine peptide 3 such as peptide 3(1-12)[MCP-4] (e.g., SEQ ID NO:65), peptide 3(1-12)[MCP-3](e.g., SEQ ID NO:66), peptide 3(1-12)[MCP-2] (e.g., SEQ ID NO:67), peptide 10 3(1-12)[eotaxin] (e.g., SEQ ID NO:68), peptide 3(1-12)[MIP1 α],(e.g., SEQ ID NO:42), peptide 3(1-12)[MIP1 β] (e.g., SEQ ID NO:43), peptide 3(1-12)[RANTES](e.g., SEQ ID NO:44), or a fragment thereof.

Another preferred embodiment of the invention includes a CXC chemokine peptide 3, a variant, an analog or a derivative thereof. A preferred CXC peptide 3 of 15 the invention is a compound of formula (III):



wherein X^2 is E or K, wherein X^3 is I, A, R or L, wherein X is D or N, wherein X^7 is Q, P or L, wherein X^4 is E, K, D, A or Q, wherein Z is A, R, S or E, wherein X^5 is P, N or K, wherein X^8 is F, W, R, I, M, L or A, wherein X^1 is L, V, Y or I, wherein 20 X^6 is K or Q, wherein a is 0-6, wherein b is 0-6, and wherein c is 1-6, with the proviso that a and b cannot both be 0.

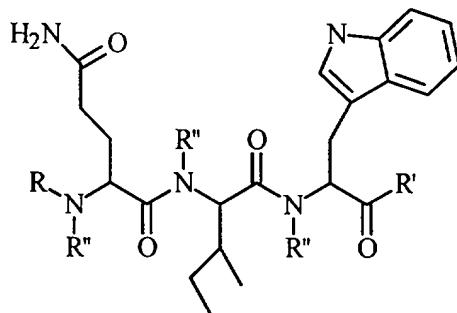
Further preferred embodiments of the invention are a CXC chemokine peptide 3 such as peptide 3(1-12)[IL8] (e.g., SEQ ID NO:40), peptide 3(1-12)[SDF-1](e.g., SEQ ID NO:38), peptide 3(1-12)[ENA-78](e.g., SEQ ID NO:41), peptide 25 3(1-12)[GRO α](e.g., SEQ ID NO:72), peptide 3(1-12)[GRO β](e.g., SEQ ID NO:73), peptide 3(1-12)[GRO γ](e.g., SEQ ID NO:74), or fragments thereof.

Yet other preferred embodiments of the invention are a CX_2C , CX_3C or C chemokine peptide 3, a variant, an analog or a derivative thereof.

Preferably, a chemokine peptide 3, its variants, analogs or derivatives inhibits the arachidonic acid pathway, e.g., inhibits the synthesis or stability, or binding, of thromboxane, prostaglandin, leukotriene, or any combination thereof.

Other compounds of the invention include compounds of formula (VIII):

5



wherein

R is (C₁-C₆)alkyl, (C₁-C₆)alkanoyl, aryl, heteroaryl, (C₁-C₆)alkoxycarbonyl, or benzyloxycarbonyl, wherein aryl, heteroaryl, and the phenyl ring of the

10 benzyloxycarbonyl can optionally be substituted with one or more (e.g. 1, 2, 3, or 4), halo, hydroxy, cyano, nitro, trifluoromethyl, trifluoromethoxy, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, (C₁-C₆)alkanoyl, (C₂-C₆)alkanoyloxy or (C₁-C₆)alkoxycarbonyl;

R' is (C₁-C₆)alkoxy, aryloxy, or NR_aR_b, wherein R_a and R_b are each independently hydrogen, (C₁-C₆)alkyl, aryl, benzyl, or phenethyl; or R_a, and R_b

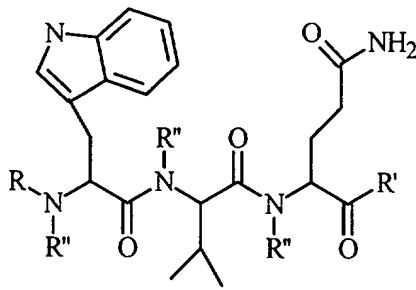
15 together with the nitrogen to which they are attached are a 5-6 membered heterocyclic ring (e.g. pyrrolidino, piperidino, or morpholino); and

each R'' is independently hydrogen, (C₁-C₆)alkyl, phenyl, benzyl, or phenethyl;

or a pharmaceutically acceptable salt thereof. Preferably, R is

20 benzyloxycarbonyl and R' is dimethylamino or diethylamino, or R is benzyloxycarbonyl; and R' is benzyloxy.

Other compounds of the invention include compounds of formula (IX):



wherein

R is (C₁-C₆)alkyl, (C₁-C₆)alkanoyl, aryl, heteroaryl, (C₁-C₆)alkoxycarbonyl, or benzyloxycarbonyl, wherein aryl, heteroaryl, and the phenyl ring of the benzyloxycarbonyl can optionally be substituted with one or more (e.g. 1, 2, 3, or 4),

5 halo, hydroxy, cyano nitro, trifluoromethyl, trifluoromethoxy, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, (C₁-C₆)alkanoyl, (C₂-C₆)alkanoyloxy or (C₁-C₆)alkoxycarbonyl;

R' is (C₁-C₆)alkoxy, aryloxy, or NR_aR_b, wherein R_a and R_b are each independently hydrogen, (C₁-C₆)alkyl, aryl, benzyl, or phenethyl; or R_a and R_b together with the nitrogen to which they are attached are a 5-6 membered

10 heterocyclic ring (e.g. pyrrolidino, piperidino, or morpholino); and

each R'' is independently hydrogen, (C₁-C₆)alkyl, phenyl, benzyl, or phenethyl;

or a pharmaceutically acceptable salt thereof. Preferably, R is benzyloxycarbonyl and R' is dimethylamino or diethylamino, or R is

15 benzyloxycarbonyl; and R' is benzyloxy.

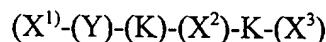
Another preferred embodiment of the invention includes a chemokine peptide 3 that is at least a tripeptide, a variant thereof or a derivative thereof. A preferred embodiment of the invention is the MCP-1 tripeptide KQK (i.e., peptide 3(9-12)[MCP-1], which specifically inhibits MCP-1, but not MIP1 α , IL8 and

20 SDF1 α , chemokine-induced activity. Other preferred embodiments of the invention include isolated and purified chemokine tripeptides that specifically inhibit IL8, MIP1 α , SDF1, murine MCP-1, MCP-2, MCP-3, and MIP1 β , e.g., KEN, SEE, KLK, KKE, KER, TQK, and SES, respectively. A further preferred embodiment of the

invention is a chemokine peptide 3 tripeptide that inhibits the activity of more than one chemokine, e.g., WVQ or WIQ. Preferably, a tripeptide of the invention is not RFK.

Yet another embodiment of the invention is a peptide which includes the 5 amino acid sequence KXK, wherein X is an amino acid, preferably one of the twenty naturally occurring amino acids, and which peptide is a chemokine antagonist, activates TGF-beta (TGF-beta1, TGF-beta2, TGF-beta3, or a combination thereof), or a combination thereof. Preferably, the peptide increases the activation of TGF-beta1. It is preferred that a peptide which includes the amino acid 10 sequence KXK is less than about 15, preferably about 10, and more preferably about 8 amino acid residues in length. Preferably, the peptide is not KFK or RPK. A further embodiment of the invention is a peptide which includes a basic amino acid residue followed by phenylalanine followed by another basic residue, wherein the peptide is not RFK, is not KFK, or does not contain RFK or KFK.

15 Another preferred peptide of the invention is a compound of formula (VII):



wherein X^2 is V, A, D, E, P, R, C, H, M, F, K, L, N, Q, Y, or I; wherein Y is absent or is an amino acid that is not R or K; and wherein X^1 and X^3 are independently 0-20 amino acid residues or absent. Preferably, X^2 is F, K, L, N, Q, Y, or I. More 20 preferably, X^2 is F, K, L, N, Q, Y, or I, and Y, X^1 and X^3 are absent.

To identify a peptide of the invention useful in the methods of the invention, a sequence comparison of chemokines from different species is performed. Then the cross-reactivity of a non-human chemokine for the human receptor is assessed. A preferred chemokine is one from a species which has the least sequence homology 25 to the corresponding human chemokine, but which still cross-reacts by binding to the human receptor. Regions which have a high degree of sequence similarity or identity are employed to prepare a peptide of the invention. For example, to identify peptides of TGF-beta having antagonist, agonist or neutral properties, the amino acid sequence of human TGF-beta was compared to that of *Xenopus*.

Peptides identified by this method include LYIDFRQDLGWKW (“T1”; SEQ ID NO:111); HEPKGYHANFC (“T2”; SEQ ID NO:112); VYYVGRK (“T3”; SEQ ID NO:113) and KVEQLSNMVVKSC (“T4”; SEQ ID NO:114). Biotinylated T1 bound to the TGF-beta receptor of THP-1 cells with an ED₅₀ of 18 μM and is a 5 receptor neutral agent (i.e., neither agonist nor antagonist). Biotinylated T2 bound to the TGF-beta receptor of THP-1 cells with an ED₅₀ of 30 μM and is a weak receptor antagonist.

Another preferred embodiment of the invention is an isolated and purified CC chemokine peptide 2, such as a peptide corresponding to SEQ ID NO:3 10 (“peptide 2(1-15)[MCP-1]”), SEQ ID NO:5 (“peptide 2(1-14)[MIP1 α]”), a fragment, a variant, an analog, or a derivative thereof. It is contemplated that chemokine peptide 2, a variant, an analog or a derivative thereof is a chemokine receptor agonist, although these therapeutic agents may exert their effect by a different mechanism, or by more than one mechanism. It is also envisioned that 15 chemokine peptide 2, a variant, an analog or a derivative thereof is a chemokine receptor antagonist. Preferably, a variant, an analog or a derivative of peptide 2 has reduced Duffy antigen binding, and also preferably, enhanced receptor binding properties, relative to the corresponding peptide 2 having a native or wild-type amino acid sequence.

20 Other preferred CC chemokine peptides 2 include peptide 2(1-14)[MIP1 β] (e.g., SEQ ID NO:60), peptide 2(1-15)[RANTES] (e.g., SEQ ID NO:61), peptide 2(1-15)[MCP-2] (e.g., SEQ ID NO:62), peptide 2(1-15)[MCP-3] (e.g., SEQ ID NO:63), peptide 2(1-15)[MCP-4] (e.g., SEQ ID NO:64), peptide 2(1-15)[eotaxin] (e.g., SEQ ID NO:75), or a fragment thereof.

25 Another preferred embodiment of the invention includes a CXC chemokine peptide 2, a variant, an analog or a derivative thereof. Preferred CC chemokine peptide 2 includes peptide 2(1-15)[IL8] (e.g., SEQ ID NO:6), peptide 2(1-15)[SDF1] (e.g., SEQ ID NO:4), peptide 2(1-15)[ENA78] (e.g., SEQ ID NO:59), peptide 2(1-

15)[GRO α] (e.g., SEQ ID NO:69), peptide 2(1-15)[GRO β] (e.g., SEQ ID NO:70), peptide 2(1-15)[GRO γ] (e.g., SEQ ID NO:71), or a fragment thereof.

Yet another preferred embodiment of the invention is a CX₂C, CX₃C or C chemokine peptide 2, a variant, an analog or a derivative thereof.

5 Other preferred peptide 2s include TSSKC (peptide 2(1-5)[MIP-1]; SEQ ID NO:102); DYFETSSQC (peptide 2(1-9)[MIP1 α]; SEQ ID NO:103); CSKPGV (peptide 2(9-14)[MIP1 α]; SEQ ID NO:104); HLKILNTPNCALQIV (peptide 2(1-5)[MIP-1 α]; SEQ ID NO:105); SYRRITSSK (peptide 2(1-5)[MCP-1]; SEQ ID NO:106); CPKEAV (peptide 2(10-15)[MCP-1]; SEQ ID NO:107); SYRRI (peptide 10 2(1-5)[MCP-1]; SEQ ID NO:108); and CSYRRITSSKSPKEAVC (SEQ ID NO:110); as well as a peptide having D isomers of the sequence VGPKSCQSSTEFYD (corresponding to residues 1-14 of peptide 2(1-14)[MIP1 α], lowercase letters are used herein to indicate D isomers as well as the letter "D" in CRD and LRD); a peptide corresponding to vaekpcksstirry; and a variant peptide 15 2 of vgpkscqsstefyd (LRD peptide 2(1-14)[MIP1 α] which includes a D isomer of serine at position 10, and the D isomer of cysteine at the amino and carboxy termini of the peptide (designated LRD-Cys₀Ser₁₀Cys₁₆ peptide 2(1-15)[MCP-1], where L = linear, F = forward, D = D isomer).

A more preferred peptide 2 of the invention is a compound of formula (XII):

20 C-(X)-(Y)-(X²)-(X⁴)-(Z)-T-(X⁵)-(X⁶)-(X¹)-C-(X⁸)-(X⁷)-(X⁹)-(X¹⁰)-V-C wherein X is S or D, wherein X² is R, K, F or Y, wherein X⁴ is R, E or F, wherein Z is T or a peptide bond, wherein X⁵ is S or N, wherein X⁶ is S or I, wherein X¹ is K, R, Q or L, wherein X⁸ is P or S, wherein X⁷ is K, R or Q, wherein X⁹ is E or P, and wherein X¹⁰ is A or G.

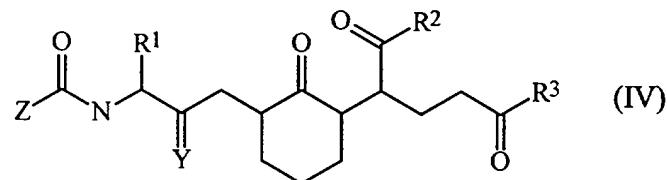
25 Also provided is an isolated and purified chemokine peptide variant, or a derivative thereof. A chemokine peptide variant has at least about 50%, preferably at least about 80%, and more preferably at least about 90% but less than 100%, contiguous amino acid sequence homology or identity to the amino acid sequence of the corresponding native chemokine, e.g., Ser, peptide 3(1-12)[MCP1] (SEQ ID

NO:11) has less than 100% contiguous homology to the corresponding amino acid sequence of MCP-1, i.e., a peptide having SEQ ID NO:1. A preferred peptide 3 variant is Leu₄Ile₁₁peptide 3(3-12)[MCP-1].

The invention also provides derivatives of chemokine peptides and peptide variants. A preferred derivative is a cyclic reverse sequence D isomer (CRD) derivative of a chemokine peptide, a variant or an analog thereof of the invention. For example, LRD derivatives of peptide 2, CRD-Cys₀Ser₁₀Cys₁₆ peptide 2[MCP-1] and CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1] are compounds of the invention that are particularly useful in the practice of the methods of the invention, as described hereinbelow.

Also provided are certain analogs of chemokines. In particular, analogs of chemokine peptide 2, chemokine peptide 3, or variants thereof are contemplated. A preferred analog of chemokine peptide 3 is an analog of WIQ. Thus, a preferred chemokine analog of the invention includes a compound of formula (IV):

15



wherein R¹ is aryl, heteroaryl, aryl(C₁-C₃)alkyl, heteroaryl(C₁-C₃)alkyl, coumaryl, coumaryl(C₁-C₃)alkyl, chromanyl or chromanyl(C₁-C₃)alkyl; wherein any aryl or heteroaryl group, or the benz-ring of any coumaryl or chromanyl group may 20 optionally be substituted with one, two or three substituents selected from the group consisting of halo, nitro, cyano, hydroxy, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, (C₁-C₆)alkanoyl, (C₂-C₆)alkanoyloxy, -C(=O)(C₁-C₆)alkoxy, C(=O)NR^gR^h, NRⁱR^j; wherein R² is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or 25 N(R^a)(R^b);

wherein R³ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or N(R^c)(R^d);

wherein Y is oxo or thioxo;

5 wherein Z is $(C_1-C_{10})alkyl$, $(C_3-C_6)cycloalkyl$, $(C_3-C_6)cycloalkyl(C_1-C_6)alkyl$,
 $(C_1-C_{10})alkoxy$, $(C_3-C_6)cycloalkyl(C_1-C_6)alkoxy$ or $N(R^e)(R^f)$; and
wherein R^a-R^j are each independently hydrogen, $(C_1-C_{10})alkyl$,
 $(C_1-C_{10})alkanoyl$, phenyl, benzyl, or phenethyl; or R^a and R^b , R^c and R^d , R^e and R^f , R^g
and R^h , or R^i and R^j together with the nitrogen to which they are attached form a ring
0 selected from pyrrolidino, piperidino, or morpholino; or a pharmaceutically
acceptable salt thereof.

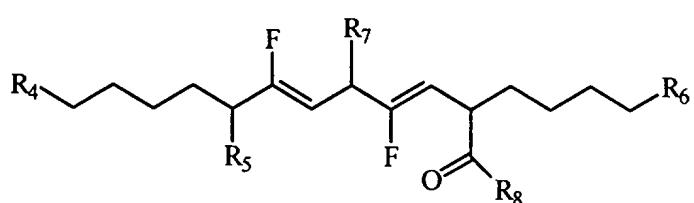
A preferred embodiment of a compound of formula (IV) includes a compound of a formula (IV) wherein R¹ is aryl, heteroaryl, coumaryl, or chromanyl.

Preferably aryl is phenyl; and heteroaryl is indolyl or pyridinyl. Another preferred

15 embodiment of a compound of formula (IV) includes a compound of a formula (IV) wherein R^2 is $N(R^a)(R^b)$; and R^3 is $N(R^c)(R^d)$. Yet another preferred embodiment of a compound of formula (IV) includes a compound of a formula (IV) wherein Z is $(C_1-C_{10})alkyl$.

A further preferred compound is a compound of formula (IV) wherein R¹ is indolyl; R² is N(R^a)(R^b); R³ is N(R^c)(R^d); Y is S; Z is hydrogen; and R^a, R^b, R^c, and R^d are each methyl.

Another preferred analog of chemokine peptide 3 is an analog of KXK. Thus, the invention includes a compound of formula (V):



wherein

R^4 is NR_kR_l ;

R^5 is NR_mR_n ;

R^6 is NR_oR_p ;

5 R^7 is the side chain of a natural or unnatural amino acid or is

$-(CH_2)_2C(=O)NR_qR_r$;

8 R^8 is hydrogen, hydroxy, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, $(C_3-$

$C_6)$ cycloalkyl(C_1-C_6)alkyl, (C_1-C_{10}) alkoxy, (C_3-C_6) cycloalkyl(C_1-C_6)alkoxy, NR_sR_t ,

the amino terminus of an amino acid or the N-terminal residue of a peptide of 2 to

10 about 25 amino acid residues;

R_k , R_l , R_o , and R_p are each independently hydrogen, (C_1-C_{10}) alkyl, $(C_3-$
 $C_6)$ cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, (C_1-C_{10}) alkanoyl, phenyl, benzyl or
phenethyl;

R_m are R_n are each independently hydrogen, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl,
15 (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, (C_1-C_{10}) alkoxy, (C_1-C_{10}) alkanoyl, $(C_1-$
 $C_{10})$ alkoxycarbonyl, 9-fluorenylmethoxycarbonyl, phenyl, benzyl, phenethyl, the C-
terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues;

R_q are R_r are each independently hydrogen, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl,
 (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, phenyl, benzyl or phenethyl;

20 wherein R_s are R_t are each independently hydrogen, (C_1-C_{10}) alkyl, $(C_3-$
 $C_6)$ cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, phenyl, benzyl or phenethyl; or a
pharmaceutically acceptable salt thereof.

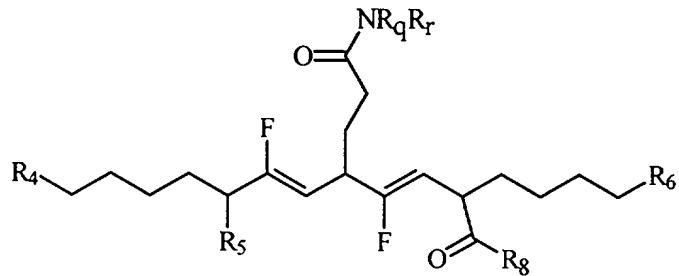
25 Preferably R_k , R_l , R_o , and R_p are each hydrogen; R_m are R_n are each
independently hydrogen, acetyl, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, propoxy, butoxy,
tert-butoxycarbonyl, 9-fluorenylmethoxycarbonyl or the C-terminal residue of an
amino acid or a peptide of 2 to about 25 amino acid residues; and R_q are R_r are each
independently hydrogen, (C_1-C_{10}) alkyl, or (C_3-C_6) cycloalkyl.

Preferably, R^7 is $-(CH_2)_2C(=O)NR_qR_r$.

Preferably, R⁷ is methyl, 3-guanidinopropyl, aminocarbonylmethyl, carboxymethyl, mercaptomethyl, (2-carboxy-2-aminoethyl)dithiomethyl, 2-carboxyethyl, 2-(aminocarbonyl)ethyl, hydrogen, 5-imadazoylmethyl, 4-amino-3-hydroxypropyl, 2-butyl, 2-methylprop-1-yl, 4-aminobutyl, 2-(methylthio)ethyl, 5 benzyl, hydroxymethyl, 1-hydroxyethyl, 3-indolylmethyl, 4-hydroxybenzyl, or isopropyl.

More preferably, R⁷ is hydrogen, benzyl, 4-hydroxybenzyl, methyl, 2-hydroxymethyl, or mercaptomethyl.

A preferred compound of formula (V) includes an analog of KGK, KFK, 10 KYK, KAK, KSK, KCK or KQK. For example, an analog of KQK includes a compound of formula (V):



wherein R⁴ is NR_kR_l;

wherein R⁵ is NR_mR_n;

15 wherein R⁶ is NR_oR_p;

wherein R⁷ is NR_qR_r;

wherein R⁸ is hydrogen, hydroxy, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy, NR_sR_t, the amino terminus of an amino acid or the N-terminal residue of a peptide of 2 to 20 about 25 amino acid residues;

wherein R_k, R_l, R_o, and R_p are each independently hydrogen, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkanoyl, phenyl, benzyl or phenethyl;

wherein R_m are R_n are each independently hydrogen, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, (C_1-C_{10}) alkoxy, (C_1-C_{10}) alkanoyl, (C_1-C_{10}) alkoxycarbonyl, 9-fluorenylmethoxycarbonyl, phenyl, benzyl, phenethyl, the C-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues;

5 wherein R_q are R_r are each independently hydrogen, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, phenyl, benzyl or phenethyl;

 wherein R_s are R_t are each independently hydrogen, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, phenyl, benzyl or phenethyl; or a pharmaceutically acceptable salt thereof.

10 Preferably R_k , R_l , R_o , and R_p are each hydrogen; R_m are R_n are each independently hydrogen, acetyl, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, propoxy, butoxy, *tert*-butoxycarbonyl, 9-fluorenylmethoxycarbonyl or the C-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues; and R_q are R_r are each independently hydrogen, (C_1-C_{10}) alkyl, or (C_3-C_6) cycloalkyl.

15 Another preferred analog of chemokine peptide 3 is an analog of WVQ (see Figure 8). Thus, the invention provides a compound of formula (VI):

(VI)

wherein

R^{10} is NR^iR^j ;

R^{11} is aryl, heteroaryl, aryl(C_1-C_3)alkyl, heteroaryl(C_1-C_3)alkyl, coumaryl, 20 coumaryl(C_1-C_3)alkyl, chromanyl or chromanyl(C_1-C_3)alkyl; wherein any aryl or heteroaryl group, or the benz-ring of any coumaryl or chromanyl group may optionally be substituted with one, two or three substituents selected from the group consisting of halo, nitro, cyano, hydroxy, (C_1-C_6) alkyl, (C_1-C_6) alkoxy, (C_1-C_6) alkanoyl, (C_2-C_6) alkanoyloxy, $-C(=O)(C_1-C_6)$ alkoxy, $C(=O)NR^eR^h$, NR^eR^f ;

25 R^{12} is (C_1-C_6) alkyl;

R¹³ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy, hydroxy, or N(R^a)(R^b);

R¹⁴ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or N(R^c)(R^d);

5 Y is oxo or thioxo;

wherein R^a-R^j are each independently hydrogen, (C₁-C₁₀)alkyl, (C₁-C₁₀)alkanoyl, phenyl, benzyl, or phenethyl; or R^a and R^b, R^c and R^d, R^e and R^f, R^g and R^h or Rⁱ and R^j together with the nitrogen to which they are attached form a ring selected from pyrrolidino, piperidino, or morpholino; or a pharmaceutically acceptable salt thereof.

10 Preferably, R¹⁰ is amino; R¹¹ is 2-benzimidazolyl; R¹² is (C₁-C₆)alkyl; R¹³ is hydroxy; and R¹⁴ is amino.

15 It is envisioned that the therapeutic agents of the invention include compounds having a chiral center that can be isolated in optically active and racemic forms.

Further provided are isolated and purified nucleic acid molecules, e.g., DNA molecules, comprising a preselected nucleic acid segment which encodes at least a portion of a chemokine, i.e., they encode a chemokine peptide or a variant thereof as described herein, e.g., a chemokine 3 peptide, a variant or derivative thereof or a 20 chemokine peptide 2, a variant or derivative thereof. For example, the invention provides an expression cassette comprising a preselected DNA segment which codes for an RNA molecule which is substantially identical (sense) to all or a portion of a messenger RNA (“target” mRNA), i.e., an endogenous or “native” chemokine mRNA. The preselected DNA segment in the expression cassette is operably linked 25 to a promoter. As used herein, “substantially identical” in sequence means that two nucleic acid sequences have at least about 65%, preferably about 70%, more preferably about 90%, and even more preferably about 98%, contiguous nucleotide sequence identity to each other. Preferably, the preselected DNA segment hybridizes under hybridization conditions, preferably under stringent hybridization

conditions, to a nucleic acid molecule encoding the corresponding native chemokine.

The present invention also provides isolated and purified DNA molecules which provide "anti-sense" mRNA transcripts of the DNA segments that encode a chemokine which, when expressed from an expression cassette in a host cell, can alter chemokine expression. As used herein, the term "antisense" means a sequence of nucleic acid which is the reverse complement of at least a portion of a RNA molecule that codes for a chemokine. Preferably, the antisense sequences of the invention are substantially complementary to a DNA segment encoding a peptide or peptide variant of the invention. As used herein, "substantially complementary" means that two nucleic acid sequences have at least about 65%, preferably about 70%, more preferably about 90%, and even more preferably about 98%, contiguous nucleotide sequence complementarity to each other. A substantially complementary RNA molecule is one that has sufficient sequence complementarity to the mRNA encoding a chemokine so as to result in a reduction or inhibition of the translation of the mRNA. It is envisioned that the duplex formed by the antisense sequence and the mRNA inhibits translation of the mRNA, as well as promotes RNA degradation, although anti-sense sequences may exert their effect by other mechanisms, or by a combination of mechanisms. Preferably, the preselected antisense DNA segment hybridizes under hybridization conditions, preferably under stringent hybridization conditions, to a nucleic acid molecule comprising the corresponding chemokine gene.

The introduction of chemokine sense or antisense nucleic acid into a cell *ex vivo* or *in vivo* can result in a molecular genetic-based therapy directed to controlling the expression of the chemokine. Thus, the introduced nucleic acid may be useful to correct or supplement the expression of the gene in patients with a chemokine-associated indication. For example, the administration of an expression vector encoding a peptide of the invention which is a chemokine receptor agonist may increase the chemokine signaling and thus be efficacious for diseases which are

characterized by decreased levels of the chemokine. Likewise, the administration of an expression vector comprising antisense chemokine sequences may be useful to prevent or treat a disorder associated with increased chemokine expression. For example, an expression vector containing antisense peptide 3(1-12)[MCP-1] which

5 is introduced into the lungs may be efficacious to prevent or treat asthma.

Also provided are pharmaceutical compositions, delivery systems, and kits comprising the therapeutic agents of the invention.

The invention further provides methods to treat chemokine-associated indications. For example, the invention provides a method of preventing or

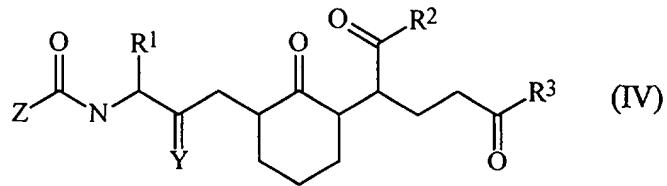
10 inhibiting an indication associated with chemokine-induced activity. The method comprises administering to a mammal afflicted with, or at risk of, the indication an amount of a chemokine peptide 3, a chemokine peptide 2, a fragment thereof, a variant thereof, a derivative thereof, a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a

15 compound of formula (XI), a compound of formula (XII), or a combination thereof, effective to prevent or inhibit said activity. Preferably, the peptide is not an IL-8 peptide, a NAP-2 peptide, or a PF4 peptide. Preferably, the administration is effective to inhibit the activity of more than one chemokine (i.e., the peptide is a pan-selective inhibitor). Preferred pan-chemokine inhibitors are WVQ, WIQ,

20 Leu₄Ile₁₁peptide 3(3-12)[MCP-1], Leu₄Ile₁₁peptide 3(1-12)[MCP-1] and CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12). These agents are useful to treat indications such as multiple sclerosis, asthma, psoriasis, allergy, rheumatoid arthritis, organ transplant rejection, and autoimmune disorders. Preferred chemokine peptides useful to treat or inhibit these indications include peptide 2 and/or peptide 3 from MCP-1, MCP-2,

25 MCP-3, MCP-4, RANTES, MIP1 α , ENA78, MIG, GRO β , eotaxin, IP10, MIP β and SDF-1.

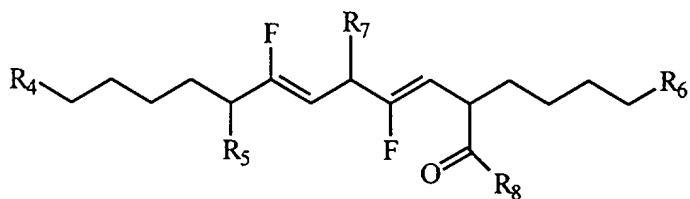
The invention also provides a method of treating a mammal afflicted with, or at risk of, an indication associated with chemokine-induced activity, comprising: administering to the mammal an effective amount of a compound of formula (IV):



wherein R¹ is aryl, heteroaryl, coumaryl or chromanyl; wherein R² is N(R^a)(R^b); wherein R³ is N(R^c)(R^d); wherein Y is oxo or thioxo; wherein Z is (C₁-C₁₀)alkyl; wherein R^a-R^d are each independently hydrogen, (C₁-C₁₀)alkyl, (C₁-C₁₀)alkanoyl, phenyl, benzyl or phenethyl; or wherein R^a and R^b, or R^c and R^d, together with the 5 nitrogen to which they are attached form a pyrrolidino, piperidino or morpholino ring; or a pharmaceutically acceptable salt thereof.

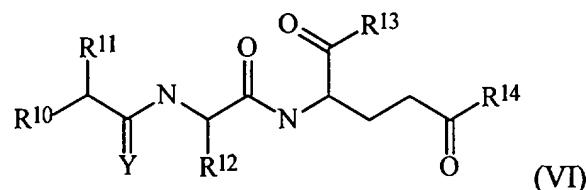
Further provided is a method of treating a mammal afflicted with, or at risk of, an indication associated with chemokine-induced activity, comprising: administering to the mammal an effective amount of a compound of formula (V):

10



C_{10})alkoxycarbonyl, 9-fluorenylmethoxycarbonyl, phenyl, benzyl, phenethyl, the C-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues; R_q are R_t are each independently hydrogen, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, phenyl, benzyl or phenethyl; wherein R_s are R_t are each independently hydrogen, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, phenyl, benzyl or phenethyl; or a pharmaceutically acceptable salt thereof.

Also provided is a method of treating a mammal afflicted with, or at risk of, an indication associated with chemokine-induced activity, comprising: administering to the mammal an effective amount of a compound of formula (VI):



10 wherein

R^{10} is NR^iR^j ;

R^{11} is aryl, heteroaryl, aryl(C_1-C_3)alkyl, heteroaryl(C_1-C_3)alkyl, coumaryl, coumaryl(C_1-C_3)alkyl, chromanyl or chromanyl(C_1-C_3)alkyl; wherein any aryl or heteroaryl group, or the benz-ring of any coumaryl or chromanyl group may 15 optionally be substituted with one, two or three substituents selected from the group consisting of halo, nitro, cyano, hydroxy, (C_1-C_6) alkyl, (C_1-C_6) alkoxy, (C_1-C_6) alkanoyl, (C_2-C_6) alkanoyloxy, $-C(=O)(C_1-C_6)$ alkoxy, $C(=O)NR^gR^h$, NR^eR^f ;

R^{12} is (C_1-C_6) alkyl;

R^{13} is (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, (C_1-C_{10}) alkoxy, (C_3-C_6) cycloalkyl(C_1-C_6)alkoxy, hydroxy, or $N(R^a)(R^b)$;

R^{14} is (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, (C_1-C_{10}) alkoxy, (C_3-C_6) cycloalkyl(C_1-C_6)alkoxy or $N(R^c)(R^d)$;

Y is oxo or thioxo;

wherein R^a-R^j are each independently hydrogen, (C_1-C_{10}) alkyl, 25 (C_1-C_{10}) alkanoyl, phenyl, benzyl, or phenethyl; or R^a and R^b , R^c and R^d , R^e and R^f , R^g

and R^h or Rⁱ and R^j together with the nitrogen to which they are attached form a ring selected from pyrrolidino, piperidino, or morpholino; or a pharmaceutically acceptable salt thereof.

The invention further provides a method to increase, augment or enhance a chemokine-associated inflammatory response in a mammal, comprising: administering to the mammal an amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof, effective to increase, augment or enhance said response. Moreover, as peptide 3, its variants and derivatives may decrease Th2 responses and increase Th1 responses, these compounds may be particularly useful to treat or prevent specific diseases in which a decrease in Th2 response and an increase in Th1 response is indicated. It is preferred that the agent employed to increase, augment or enhance the chemokine-associated inflammatory response is not YNFTNRKISVQRLASYRRITSSK. These therapeutic agents are useful to increase an inflammatory response to, for example, intracellular pathogens or parasites, which often are associated with a poor immune response. Thus, these agents may be useful to treat or prevent tuberculosis and malaria. Therefore, the invention also provides a therapeutic method to prevent or treat parasitic infection.

The invention also provides a method of preventing or inhibiting an indication associated with histamine release from basophils or mast cells. The method comprises administering to a mammal at risk of, or afflicted with, the indication an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula

(X), a compound of formula (XI), a compound of formula (XII), or a combination thereof.

Also provided is a method of preventing or inhibiting an indication associated with monocyte, macrophage, neutrophil, B cell, T cell or eosinophil recruitment, or B cell or T cell activation or proliferation. The method comprises administering an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. For example, a chemokine peptide 3, a chemokine peptide 2, a variant thereof, or a derivative thereof may be useful to prevent or treat autoimmune or granulomatous indications.

Further provided is a therapeutic method to prevent or treat vascular indications, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof, wherein the indication is coronary artery disease, myocardial infarction, unstable angina pectoris, atherosclerosis or vasculitis. Preferred chemokine peptides for this embodiment of the invention include chemokine peptides of MCP-1, RANTES, GRO α , MIP1 α , IP10, MCP-4, and MIP1 β .

The invention also provides a method to prevent or treat an autoimmune disorder. The method comprises administering to a mammal in need of said therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a

compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. A preferred variant of peptide 3 useful to prevent or treat autoimmune disorders is Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14) or peptide 3 having
5 WVQ. A preferred chemokine peptide 3 for use in preventing or treating multiple sclerosis includes SEE and peptide 3(1-14)[MIP1 α] (SEQ ID NO:42). Other preferred peptides are chemokine peptides of RANTES.

Further provided is a method to modulate the chemokine-induced activity of macrophage, B cells, T cells or other hematopoietic cells, e.g., neutrophils,
10 eosinophils or mast cells, at a preselected physiological site. The method comprises administering a dosage form comprising an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a
15 compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof, wherein the dosage form is linked, either covalently or noncovalently, to a targeting moiety. The targeting moiety binds to a cellular component at the preselected physiological site.

Moreover, it is also envisioned that an agent of the invention may be a
20 targeting moiety, as some of the agents are selective chemokine inhibitors, rather than pan-chemokine inhibitors. For example, an agent of the invention, e.g., peptide 2, may be useful in the targeted delivery of an isotope or other cytotoxic molecule to red cells for the treatment of disorders such as erythroid leukemia, erythroid
myelosis, polycythemia vera or other erythroid dysplasias. Similarly, an agent of the
25 invention that specifically targets a particular cell type may be useful in diagnostics. Thus, these agents can be radiolabeled (Chianelli et al., Nucl. Med. Comm., 18, 437 (1997)), or labeled with any other detectable signal, such as those useful in diagnostic imaging (e.g., MRI and CAT) to image sites of inflammation in disorders like rheumatoid arthritis and diabetes mellitus (type I).

Also provided is a therapeutic method to augment an immune response. The method comprises administering to a mammal an immunogenic moiety and an amount of a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula 5 (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof, wherein the amount is effective to augment the immune response of the mammal to the immunogenic moiety. Thus, the invention also provides a vaccine comprising an immunogenic moiety and an 10 amount of a chemokine peptide 2, a variant thereof or a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. As used herein, an “immunogenic 15 moiety” means an isolated or purified composition or compound (e.g., a purified virus preparation or a native or recombinant viral or bacterial antigen) that, when introduced into an animal, preferably a mammal, results in a humoral and/or cellular immune response by the animal to the composition or compound. Also provided are modified vaccines, wherein the immunogenic moiety is coupled to peptide 2, a 20 variant or derivative thereof. Peptide 2 increases the binding of the modified vaccine to the red blood cell pool and blocks Duffy binding of chemokines and so prolongs the residency time of the vaccine in the circulation and decreases chemokine-induced activity, either of which result in an augmented antibody response. It is envisaged that the modified vaccine is delivered by the same routes 25 as those used for unmodified immunogen (e.g., intravenous, intramuscular or orally).

The invention also provides a therapeutic method to prevent or inhibit asthma. The method comprises administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant

thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. As described hereinbelow in Example 12, a peptide of the invention inhibited cellular inflammation and IgE responses in the lung of mice exposed to ovalbumin. Preferably, in this embodiment of the invention, a therapeutic agent is administered to the upper and/or lower respiratory tract. Preferred peptides useful in this embodiment of the invention are chemokine peptides of RANTES, MCP-1 and 10 MIP1 α .

Further provided is a therapeutic method to prevent or inhibit viral, e.g., poxvirus, herpesvirus (e.g., *Herpesvirus samiri*), cytomegalovirus (CMV) or lentivirus, infection or replication. The method comprises administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a 15 chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), or a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. Preferably, the therapeutic agents are employed to 20 prevent or treat HIV. More preferably, the agent is administered before, during or after the administration of an anti-viral agent, e.g., for HIV AZT, a non-nucleoside reverse transcriptase inhibitor, a protease inhibitor or a combination thereof. It is also envisioned that a combination of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound 25 of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), or a compound of formula (XII) may be useful in the anti-viral methods and compositions of the invention. Preferred chemokine

peptides useful to prevent or inhibit viral infection are those from IP10, MIP1 α , MIP1 β , SDF-1, IL-8, GRO α , RANTES and MCP-1.

A therapeutic method to prevent or treat low bone mineral density is also provided. The method comprises administering to a mammal in need of such

5 therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination

10 thereof. A preferred derivative of a variant of peptide 3 to prevent or treat low mineral bone density is CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1]. A preferred fragment of SEQ ID NO:1 useful in preventing or treating low mineral bone density is KQK.

Also provided is a method of suppressing tumor growth in a vertebrate

15 animal comprising administering to said vertebrate a therapeutically effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. Preferably, the method increases or enhances macrophage, B cell-, T cell- or other immune cell-associated activity at a tumor site. A preferred peptide for use in this embodiment of the invention is a MCP-1 peptide.

Further provided is a method for preventing or treating rheumatoid arthritis

25 in a mammal, comprising: administering to the mammal an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound

of formula (XII), or a combination thereof. For this embodiment of the invention, a preferred peptide is a MCP-1, MIP1 α , MIP1 β , GRO α , and ENA78 peptide.

Also provided is a method to prevent or treat organ transplant rejection. The method comprises administering an effective amount of a chemokine peptide 3, a 5 chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof.

10 Further provided is a method for preventing or treating psoriasis in a mammal, comprising: administering to the mammal an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of 15 formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. Preferred peptides to prevent or treat psoriasis are peptides of MCP-1, RANTES, MIP1 α , MIG, IP10, GRO β , GRO α or MCP-3. A preferred derivative to prevent or treat psoriasis is a CRD-derivative of peptide 3.

20 Also provided is a method to enhance wound healing. The method comprises administering an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of 25 formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof.

Further provided is a method for preventing or treating an allergy in a mammal, comprising: administering to the mammal an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof,

a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof. Preferred peptides to prevent or treat
5 allergies include peptides of RANTES, MIP1 α , MCP-1, MCP-2, MCP-3, MCP-4, eotaxin or MIP1 β .

Yet another embodiment of the invention is a method to prevent or inhibit an indication associated with elevated TNF- α . The method comprises administering an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof,
10 a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), a compound of formula (VII), a compound of formula (VIII), a compound of formula (IX), a compound of formula (X), a compound of formula (XI), a compound of formula (XII), or a combination thereof.

The invention also provides methods in which the nucleic acid molecules of
15 the invention are administered to a mammal afflicted with, or at risk of, an indication associated with a chemokine-induced activity.

The invention also provides methods whereby the pharmacokinetics of desirable pharmaceutical agents may be modulated. In particular, agents which are normally rapidly cleared from the circulation may be retained longer by the addition
20 of a peptide of the invention that has affinity for the Duffy antigen on red blood cells. This methodology may be particularly suited to modifying the pharmacokinetics of other biologically active, pharmaceutically useful peptides, as well as larger polypeptide or proteins. For example, a Duffy binding peptide (such as peptide 2[MCP-1] may be coupled or linked, either covalently or non-covalently,
25 to a molecule such as recombinant human growth hormone (HGH) or insulin, and administered via a depo injection. By partitioning the modified HGH to the red blood cells, HGH may have much more suitable pharmacokinetics, with longer half times and less rapid changes in plasma concentrations. In another example, insulin coupled to a peptide of the invention may be particularly useful as a treatment for

the highly insulin-resistant type II diabetic whose disease has progressed significantly. It is also envisioned that other small molecules may be coupled to Duffy binding molecules in a manner which preserves the intended function of the active molecule and of the Duffy binding molecule. For coupling to recombinant 5 proteins and peptides, Duffy binding peptides are preferred. For coupling to small molecule drugs, analogs (e.g., isosteres) of Duffy binding peptides are preferred.

Brief Description of the Figures

Figure 1 is a schematic of the trans-well migration assay. In most 10 experiments, the peptide (wavy line) is added to the upper well with about 50,000 cells (O). The upper and lower wells are separated by a 5 μ m or 8 μ m pore size PVP-free membrane (---). Chemokine (●) is added to the lower well. After 4 hours, the number of cells that have migrated through the membrane are measured (O in lower well).

15 Figure 2 shows a dose-response curve for the peptide 3 (SEQ ID NO:1) inhibition of MCP-1-induced THP-1 cell migration.

Figure 3 shows the reverse transcriptase activity present in the culture medium at day 21 after infection of Jurkat cells with a T-tropic HIV. Peptides were added on day 0, one hour prior to infection of the cells with HIV isolate. The full 20 length chemokine SDF-1 α was used as a positive control.

Figure 4 shows the structure of CRD-Cys₁₃leu₄ile₁₁peptide 3[MCP-1](3-12)[MCP-1], which is cyclized via disulphide bonds. The main chain α carbons are indicated by C_D which indicates that the D form of the amino acid is present.

Figure 5 depicts a schematic of inhibition of cell migration via binding of a 25 therapeutic agent of the invention to a chemokine receptor. C^RC = a therapeutic agent of the invention. Chemokine receptors are shown as blackened rectangles.

Figure 6 depicts a schematic of the inhibition of HIV entry by an agent of the invention.

Figure 7 shows the dose-dependent inhibition of inflammation (A) and endotoxemia (B) in animal models by peptide 3 (CRD-Cys₁₃Leu₄Ile₁₁ peptide 3(3-12) [MCP-1] = NR58-3.14.3).

Figure 8 shows a preferred analog of peptide WVQ.

5 Figure 9 shows a graph of the number of macrophage at the site of LPS administration in a rat in the presence or absence of a peptide of the invention.

Figure 10 shows a graph of the number of B cells at the site of LPS administration in a rat in the presence or absence of a peptide of the invention.

10 Figure 11 shows a graph of the fraction of HIV infected THP-1 cells in the presence of peptide 2 or peptide 3 using a quantitative immunofluorescent (QIF) assay.

Figure 12 depicts codons for various amino acids.

Figure 13 depicts exemplary amino acid substitutions.

Figure 14 shows exemplary therapeutic agents of the invention.

15 Figure 15 shows the Duffy binding affinity and inhibition of THP-1 migration for selected peptide 2 compounds. LFL = linear forward L isomer; LRD = linear reverse D isomer; CRD = cyclic reverse D isomer; CFL = cyclic forward L isomer.

20 Figure 16 summarizes binding and ED₅₀ data for selected peptides of the invention.

Figure 17 shows an exemplary protocol to test agents in a rat dermal inflammation model (CRD-Cys₁₃Leu₄Ile₁₁ peptide 3(3-12)[MCP-1] = NR58-3.14.3).

Detailed Description of the Invention

25 Definitions

“Chemokines” refers to a family of proinflammatory signaling molecules which act on macrophage, B cells, T cells, neutrophils, eosinophils, basophils, mast cells, smooth muscle cells, e.g., vascular smooth muscle cells, and the like (e.g., by affecting their migration, proliferation, or degranulation, or the immunomodulation

of T cell development to Th1 and Th2 subtypes). Preferred chemokines are primate in origin, e.g., human, although the invention includes other mammalian chemokines, such as those of bovine, ovine, equine, canine, feline or rodent origin, as well as virally encoded chemokines. Chemokines include, but are not limited to,

5 MCP-1 (SEQ ID NO:16), MCP-2 (SEQ ID NO:17), MCP-3 (SEQ ID NO:18), MIG (SEQ ID NO:45), MIP1 α (SEQ ID NO:19), MIP1 β (SEQ ID NO:20), RANTES (SEQ ID NO:21), PF4 (SEQ ID NO:46), I-309 (SEQ ID NO:47), HCC-1 (SEQ ID NO:48), eotaxin (SEQ ID NO:25), C10 (SEQ ID NO:49), CCR-2 (SEQ ID NO:50), ENA-78 (SEQ ID NO:52), GRO α (SEQ ID NO:24), GRO β (SEQ ID NO:53), IL-8 (SEQ ID NO:23), IP-10 (SEQ ID NO:54), SDF1 α (SEQ ID NO:22), SDF1 β (SEQ ID NO:56), GRO α (SEQ ID NO:57), MIP3 α , TCA-3, CTAPIII, MARC/FYK, β -thromboglobulin, GCP-2, PBP, HC14, MDC, TECK, PARC, 6Ckine, fractaline, DC-CK1, LIX, TARC, LARC, MIG, Ck β 8, CCF18/MRP-2, CCIII, CK α 2, H1305, Dvic-1, MGSA, Ck β 4, DGWCC, TCA4, dendrokine (see WO 97/29192),

10 15 CC2/HCC1, CC3, and MIP1 τ , as well as virally encoded chemokines such as vMIP-I, vMIP-II and vMIP-III (see Kledal et al., *Science*, 277, 1656 (1997)). “CXC” or “ α ” chemokines include, but are not limited to, IL-8, PF4, IP10, NAP-2, GRO α , GRO β , GRO γ , SDF1, MIP2, MGSA, γ IP, CTAPIII, β -thromboglobulin, MIG, PBP, NAP-2 and ENA78. “CC” or “ β ” chemokines include, but are not limited to, MCP-20 1, MCP-2, MCP-3, MCP-4, MCP-5, RANTES, eotaxin, LARC, TARC, C10, MIP1 α , MIP1 β , I309, HCC-1, CK β 8, CCF18/MRP-2, MIP1 τ . A third type of chemokines are “C” chemokines, e.g., lymphotactin. A fourth type of chemokines are “CX₃C” chemokines such as fractaline or neurotactin (Rollins et al., *Blood*, 90, 404 (1997)). A fifth type of chemokines, CX₂C chemokines, include CCIII.

25 “Peptide 3” refers to a peptide derived from a chemokine, which is generally located in the carboxy-terminal half of the chemokine, and which inhibits the activity of at least the corresponding native chemokine, as determined by methods well known to the art. Peptide 3 comprises no more than 30, preferably about 3 to about 25, more preferably about 3 to about 15, and even more preferably about 3 to

about 11, peptidyl residues which have 100% contiguous amino acid sequence homology or identity to the amino acid sequence of the corresponding native chemokine, preferably a mammal chemokine, e.g., a primate chemokine such as a human chemokine, or a virally-encoded chemokine. For example, a preferred 5 peptide 3 of MCP-1 that inhibits at least the activity of MCP-1 is peptide 3(1-12)[MCP-1], e.g., a peptide which has an amino acid sequence corresponding to SEQ ID NO:1, or a fragment or derivative thereof. Another preferred embodiment of the invention is peptide 3(3-12)[MCP-1], e.g., a peptide having an amino acid sequence corresponding to SEQ ID NO:7, or a fragment or derivative thereof.

10 Preferably, a chemokine peptide 3 of the invention does not include a peptide of IL-8, PF-4 or NAP-2.

An alignment of chemokine amino acid sequences, such as the alignment depicted in Table 1, provides a general method to identify the location of peptide 3 sequences in chemokines. Generally, peptide 3 in non-MCP-1 chemokines 15 corresponds to about residue 46 to about residue 67 of mature human MCP-1. Moreover, it is envisioned that peptide 3 may comprise moieties other than the amino acid sequence which inhibits chemokine activity, e.g., amino acid residues not present in the native chemokine (i.e., a fusion protein), nucleic acid molecules or targeting moieties such as antibodies or fragments thereof or biotin, so long as these 20 moieties do not substantially reduce the biological activity of peptide 3. A substantial reduction in activity means a reduction in activity of greater than about 99%.

“Peptide 2” refers to a peptide derived from a chemokine, which is generally located in the amino-terminal two-thirds of the chemokine, and which does not 25 include the amino-terminal about 20 to about 24 amino acid residues of the native mature chemokine. Generally, peptide 2 is a chemokine agonist, but peptide 2 may also have neither agonist or antagonist activities (i.e., it is “neutral”), or may be a chemokine antagonist, so long as the peptide specifically binds to at least one chemokine receptor. Peptide 2 comprises no more than 30, preferably about 3 to

about 25, more preferably about 10 to about 25, and even more preferably about 10 to about 18, peptidyl residues which have 100% contiguous amino acid sequence homology or identity to the amino acid sequence of the corresponding native chemokine. For example, a preferred peptide 2 of MCP-1 is peptide 2(1-15)[MCP-1], for example a peptide which has an amino acid sequence corresponding to SEQ ID NO:3, or a fragment or derivative thereof. A more preferred peptide 2 is a peptide 2 comprising at least one D isomer. Preferably, a chemokine peptide 2 of the invention is not peptide 2[PF4], peptide 2[IL-8], peptide 2 [NAP-2] or YNFTNRKISVQLASYRRITSSK.

10 An alignment of chemokine amino acid sequences, such as the alignment depicted in Table 1, provides a general method to identify the location of peptide 2 sequences in other chemokines. Generally, peptide 2 in non-MCP-1 chemokines corresponds to about residue 27 to about residue 45 on mature human MCP-1. It is also envisioned that peptide 2 may comprise moieties other than the amino acid sequence which mimics, enhances or does not affect (i.e., neutral) chemokine activity, e.g., amino acid residues not present in the native chemokine, nucleic acid molecules or targeting moieties such as those described above for peptide 3, so long as these moieties do not substantially alter the biological activity of peptide 2. A substantial alteration in activity means an alteration of greater than about 99%.

15 20 Also preferably, a peptide, variant, analog or derivative of the invention, has increased affinity for at least one chemokine receptor, e.g., about 1 μ M to about 1 nM, more preferably about 1 nM to about 1 pM, and also preferably has decreased Duffy binding, relative to a corresponding peptide having the native (“wild-type”) sequence or relative to the corresponding native chemokine. However, certain populations have individuals who are Duffy⁻, e.g., a certain percentage of African Americans are Duffy⁻. Thus, agents useful to treat these populations may have Duffy binding affinity that is equal to or greater than that of the corresponding native chemokine.

As used herein, the terms “isolated and/or purified” refer to *in vitro* preparation, isolation and/or purification of a therapeutic agent of the invention, so that it is not associated with *in vivo* substances. Thus, with respect to an “isolated nucleic acid molecule”, which includes a polynucleotide of genomic, cDNA, or synthetic origin or some combination thereof, the “isolated nucleic acid molecule”

5 (1) is not associated with all or a portion of a polynucleotide in which the “isolated nucleic acid molecule” is found in nature, (2) is operably linked to a polynucleotide which it is not linked to in nature, or (3) does not occur in nature as part of a larger sequence. An isolated nucleic acid molecule means a polymeric form of

10 nucleotides of at least 10 bases in length, either ribonucleotides or deoxynucleotides or a modified form of either type of nucleotide. The term includes single and double stranded forms of DNA. The term “oligonucleotide” referred to herein includes naturally occurring, and modified nucleotides linked together by naturally occurring, and non-naturally occurring oligonucleotide linkages. Oligonucleotides are a

15 polynucleotide subset with 200 bases or fewer in length. Preferably, oligonucleotides are 10 to 60 bases in length and most preferably 12, 13, 14, 15, 16, 17, 18, 19, or 20 to 40 bases in length. Oligonucleotides are usually single stranded, e.g., for probes; although oligonucleotides may be double stranded, e.g., for use in the construction of a variant. Oligonucleotides of the invention can be either sense

20 or antisense oligonucleotides. The term “naturally occurring nucleotides” referred to herein includes deoxyribonucleotides and ribonucleotides. The term “modified nucleotides” referred to herein includes nucleotides with modified or substituted sugar groups and the like. The term “oligonucleotide linkages” referred to herein includes oligonucleotides linkages such as phosphorothioate, phosphorodithioate,

25 phosphoroselenoate, phosphorodiselenoate, phosphoroanilothioate, phosphoranyliladate, phosphoroamidate, and the like. An oligonucleotide can include a label for detection, if desired.

The term “isolated polypeptide” means a polypeptide encoded by cDNA or recombinant RNA, or is synthetic origin, or some combination thereof, which

isolated polypeptide (1) is not associated with proteins found in nature, (2) is free of other proteins from the same source, e.g., free of human proteins, (3) is expressed by a cell from a different species, or (4) does not occur in nature.

The term “sequence homology” means the proportion of base matches

- 5 between two nucleic acid sequences or the proportion amino acid matches between two amino acid sequences. When sequence homology is expressed as a percentage, e.g., 50%, the percentage denotes the proportion of matches over the length of sequence from a chemokine that is compared to some other sequence. Gaps (in either of the two sequences) are permitted to maximize matching; gap lengths of
- 10 15 bases or less are usually used, 6 bases or less are preferred with 2 bases or less more preferred. When using oligonucleotides as probes or treatments, the sequence homology between the target nucleic acid and the oligonucleotide sequence is generally not less than 17 target base matches out of 20 possible oligonucleotide base pair matches (85%); preferably not less than 9 matches out of 10 possible base
- 15 pair matches (90%), and more preferably not less than 19 matches out of 20 possible base pair matches (95%).

The term “selectively hybridize” means to detectably and specifically bind. Polynucleotides, oligonucleotides and fragments of the invention selectively hybridize to nucleic acid strands under hybridization and wash conditions that

- 20 minimize appreciable amounts of detectable binding to nonspecific nucleic acids. High stringency conditions can be used to achieve selective hybridization conditions as known in the art and discussed herein. Generally, the nucleic acid sequence homology between the polynucleotides, oligonucleotides, and fragments of the invention and a nucleic acid sequence of interest is at least 65%, and more typically 25 with preferably increasing homologies of at least about 70%, about 90%, about 95%, about 98%, and 100%.

Two amino acid sequences are homologous if there is a partial or complete identity between their sequences. For example, 85% homology means that 85% of the amino acids are identical when the two sequences are aligned for maximum

matching. Gaps (in either of the two sequences being matched) are allowed in maximizing matching; gap lengths of 5 or less are preferred with 2 or less being more preferred. Alternatively and preferably, two protein sequences (or polypeptide sequences derived from them of at least 30 amino acids in length) are homologous,

5 as this term is used herein, if they have an alignment score of at more than 5 (in standard deviation units) using the program ALIGN with the mutation data matrix and a gap penalty of 6 or greater. See Dayhoff, M. O., in *Atlas of Protein Sequence and Structure*, 1972, volume 5, National Biomedical Research Foundation, pp. 101-110, and Supplement 2 to this volume, pp. 1-10. The two sequences or

10 parts thereof are more preferably homologous if their amino acids are greater than or equal to 50% identical when optimally aligned using the ALIGN program.

The term "corresponds to" is used herein to mean that a polynucleotide sequence is homologous (i.e., is identical, not strictly evolutionarily related) to all or a portion of a reference polynucleotide sequence, or that a polypeptide sequence is

15 identical to a reference polypeptide sequence. In contradistinction, the term "complementary to" is used herein to mean that the complementary sequence is homologous to all or a portion of a reference polynucleotide sequence. For illustration, the nucleotide sequence "TATAC" corresponds to a reference sequence "TATAC" and is complementary to a reference sequence "GTATA".

20 The following terms are used to describe the sequence relationships between two or more polynucleotides: "reference sequence", "comparison window", "sequence identity", "percentage of sequence identity", and "substantial identity". A "reference sequence" is a defined sequence used as a basis for a sequence comparison; a reference sequence may be a subset of a larger sequence, for example,

25 as a segment of a full-length cDNA or gene sequence given in a sequence listing, or may comprise a complete cDNA or gene sequence. Generally, a reference sequence is at least 20 nucleotides in length, frequently at least 25 nucleotides in length, and often at least 50 nucleotides in length. Since two polynucleotides may each (1) comprise a sequence (i.e., a portion of the complete polynucleotide sequence) that is

similar between the two polynucleotides, and (2) may further comprise a sequence that is divergent between the two polynucleotides, sequence comparisons between two (or more) polynucleotides are typically performed by comparing sequences of the two polynucleotides over a “comparison window” to identify and compare local regions of sequence similarity.

A “comparison window”, as used herein, refers to a conceptual segment of at least 20 contiguous nucleotides and wherein the portion of the polynucleotide sequence in the comparison window may comprise additions or deletions (i.e., gaps) of 20 percent or less as compared to the reference sequence (which does not comprise additions or deletions) for optimal alignment of the two sequences.

Optimal alignment of sequences for aligning a comparison window may be conducted by the local homology algorithm of Smith and Waterman (1981) Adv. Appl. Math. 2: 482, by the homology alignment algorithm of Needleman and Wunsch (1970) J. Mol. Biol. 48: 443, by the search for similarity method of Pearson and Lipman (1988) Proc. Natl. Acad. Sci. (U.S.A.) 85: 2444, by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package Release 7.0, Genetics Computer Group, 575 Science Dr., Madison, Wis.), or by inspection, and the best alignment (i.e., resulting in the highest percentage of homology over the comparison window) generated by the various methods is selected.

The term “sequence identity” means that two polynucleotide sequences are identical (i.e., on a nucleotide-by-nucleotide basis) over the window of comparison. The term “percentage of sequence identity” means that two polynucleotide sequences are identical (i.e., on a nucleotide-by-nucleotide basis) over the window of comparison. The term “percentage of sequence identity” is calculated by comparing two optimally aligned sequences over the window of comparison, determining the number of positions at which the identical nucleic acid base (e.g., A, T, C, G, U, or I) occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the

window of comparison (i.e., the window size), and multiplying the result by 100 to yield the percentage of sequence identity. The terms “substantial identity” as used herein denote a characteristic of a polynucleotide sequence, wherein the polynucleotide comprises a sequence that has at least 85 percent sequence identity,

5 preferably at least 90 to 95 percent sequence identity, more usually at least 99 percent sequence identity as compared to a reference sequence over a comparison window of at least 20 nucleotide positions, frequently over a window of at least 20-50 nucleotides, wherein the percentage of sequence identity is calculated by comparing the reference sequence to the polynucleotide sequence which may

10 include deletions or additions which total 20 percent or less of the reference sequence over the window of comparison. The reference sequence may be a subset of a larger sequence, for example, as a segment of human MCP-1.

As applied to polypeptides, the term “substantial identity” means that two peptide sequences, when optimally aligned, such as by the programs GAP or

15 BESTFIT using default gap weights, share at least about 80 percent sequence identity, preferably at least about 90 percent sequence identity, more preferably at least about 95 percent sequence identity, and most preferably at least about 99 percent sequence identity.

As used herein, the terms “label” or “labeled” refer to incorporation of a

20 detectable marker, e.g., by incorporation of a radiolabeled amino acid or attachment to a polypeptide of biotinyl moieties that can be detected by marked avidin (e.g., streptavidin containing a fluorescent marker or enzymatic activity that can be detected by optical or colorimetric methods). Various methods of labeling polypeptides are known in the art and may be used. Examples of labels for

25 polypeptides include, but are not limited to, the following: radioisotopes (e.g., ^3H , ^{14}C , ^{35}S , ^{125}O , ^{131}I), fluorescent labels (e.g., FITC, rhodamine, lanthanide, phosphors), enzymatic labels (e.g., horseradish peroxidase, β -galactosidase, luciferase, alkaline phosphatase), chemiluminescent, biotinyl groups, predetermined polypeptide epitopes recognized by a secondary reporter (e.g., leucine zipper pair sequences,

binding sites for secondary antibodies, metal binding domains, epitope tags). In some embodiments, labels are attached by spacer arms of various lengths to reduce potential steric hindrance.

As used herein, “substantially pure” means an object species is the

5 predominant species present (i.e., on a molar basis it is more abundant than any other individual species in the composition), and preferably a substantially purified fraction is a composition wherein the object species comprises at least about 50 percent (on a molar basis) of all macromolecular species present. Generally, a substantially pure composition will comprise more than about 80 percent of all

10 macromolecular species present in the composition, more preferably more than about 85%, about 90%, about 95%, and about 99%. Most preferably, the object species is purified to essential homogeneity (contaminant species cannot be detected in the composition by conventional detection methods) wherein the composition consists essentially of a single macromolecular species.

15 An isolated “chemokine peptide variant” of peptide 3 or peptide 2 is a peptide comprising no more than 30, preferably about 3 to about 25, and more preferably about 3 to about 18, and even more preferably about 3 to about 11, peptidyl residues which have at least 50%, preferably at least about 80%, and more preferably at least about 90% but less than 100%, contiguous amino acid sequence

20 homology or identity to the amino acid sequence of the corresponding native chemokine, e.g., Ser₇ peptide 3(1-12)[MCP1] (SEQ ID NO:11) has less than 100% homology to the corresponding amino acid sequence of MCP-1, i.e., peptide 3(1-12)[MCP-1] (SEQ ID NO:1). A variant of the invention may include amino acid residues not present in the corresponding native chemokine, and internal deletions

25 relative to the corresponding native chemokine. Chemokine peptide variants include peptides having at least one D-amino acid.

Chemokine peptides or peptide variants which are subjected to chemical modifications, such as esterification, amidation, reduction, protection and the like, are referred to as chemokine “derivatives.” For example, a modification known to

improve the stability and bioavailability of peptides *in vivo* is the cyclization of the peptide, for example through one or more disulfide bonds. A preferred modification is the synthesis of a cyclic reverse sequence derivative (CRD) of a peptide of the invention. A linear peptide is synthesized with all D-form amino acids using the 5 reverse (i.e., C-terminal to N-terminal) sequence of the peptide. If necessary, additional cysteine residues are added to the N and C termini (if the peptide sequence does not already have N and C terminal cys residues), thereby allowing oxidative cyclization. However, the term “CRD” includes cyclization by other mechanisms, e.g., via a peptidyl bond, and the like. A preferred derivative of the 10 invention is CRD-Cys₀Cys₁₃Leu₄Ile₁₁peptide 3[MCP-1] or CRD-Cys₁₃Leu₄Ile₁₁ peptide 3(3-12)[MCP-1].

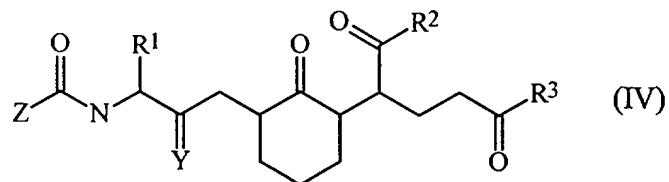
Also included within the scope of the term “derivative” is linear reverse D (LRD) and cyclized forward L (CFL) derivatives. LRD derivatives have the reverse (i.e., C-terminal to N-terminal) sequence of the peptide with all D-form amino acids, 15 but are not cyclized. CFL derivatives have the forward (i.e., N-terminal to C-terminal) sequence of the peptide with all L-form amino acids, but with additional N and C terminal cys residues (if the peptide sequence does not already have cys residues at either the N or the C terminal position), followed by oxidative cyclization, or cyclization by an alternative method. Other “derivatives” of the 20 invention include branched peptides, circular, branched and branched circular peptides.

A “chemokine analog” means a moiety that mimics or inhibits a chemokine-induced activity, or binds to or near a chemokine receptor but does not mimic or inhibit chemokine activity (neutral), wherein the portion of the moiety that mimics 25 or inhibits the chemokine-induced activity, or binds to or near the receptor but is neutral, is not a peptide, and wherein the active portion of the analog is not a nucleic acid molecule. As used herein, the term “mimics” means that the moiety induces an activity that is induced by a native chemokine, but that the induction by the analog is

not necessarily of the same magnitude as the induction of activity by the native chemokine.

It is also envisioned that the chemokine peptides, variants, analogs and derivatives thereof, of the invention may comprise moieties other than the portion 5 which inhibits or mimics chemokine activity, or binds to or near a chemokine receptor without eliciting or inhibiting signaling, e.g., peptide or polypeptide molecules, such as antibodies or fragments thereof or fusion proteins, nucleic acid molecules, sugars, lipids, fats, a detectable signal molecule such as a radioisotope, e.g., gamma emitters, paramagnetic molecules or sound wave emitters, small 10 chemicals, metals, salts, synthetic polymers, e.g., polylactide and polyglycolide, surfactants and glycosaminoglycans, which preferably are covalently attached or linked to the portion of the peptide, variant, analog or derivative that mimics or inhibits the chemokine-induced activity, so long as the other moieties do not alter the biological activity of the peptide, variant, analog or derivative. Also envisioned 15 is a chemokine peptide, variant, analog or derivative that is non-covalently associated with the moieties described above.

A preferred chemokine analog of the invention is a compound of formula (IV):



20

wherein R¹ is aryl, heteroaryl, aryl(C₁-C₃)alkyl, heteroaryl(C₁-C₃)alkyl, coumaryl, coumaryl(C₁-C₃)alkyl, chromanyl or chromanyl(C₁-C₃)alkyl; wherein any aryl or heteroaryl group, or the benz-ring of any coumaryl or chromanyl group may optionally be substituted with one, two or three substituents selected from the group

consisting of halo, nitro, cyano, hydroxy, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, (C₁-C₆)alkanoyl, (C₂-C₆)alkanoyloxy, -C(=O)(C₁-C₆)alkoxy, C(=O)NR^gR^h, NRⁱR^j;
wherein R² is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or N(R^a)(R^b);
5 wherein R³ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or N(R^c)(R^d);
wherein Y is oxo or thioxo;
wherein Z is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or N(R^e)(R^f); and
10 wherein R^a-R^j are each independently hydrogen, (C₁-C₁₀)alkyl, (C₁-C₁₀)alkanoyl, phenyl, benzyl, or phenethyl; or R^a and R^b, R^c and R^d, R^e and R^f, R^g and R^h, or Rⁱ and R^j together with the nitrogen to which they are attached form a ring selected from pyrrolidino, piperidino, or morpholino; or a pharmaceutically acceptable salt thereof.

15 A preferred embodiment of a compound of formula (IV) includes a compound of formula (IV) wherein R¹ is aryl, heteroaryl, coumaryl, or chromanyl. Preferably aryl is phenyl; and heteroaryl is indolyl or pyridinyl. Another preferred embodiment of a compound of formula (IV) includes a compound of a formula (IV) wherein R² is N(R^a)(R^b); and R³ is N(R^c)(R^d). Yet another preferred embodiment of
20 a compound of formula (IV) includes a compound of a formula (IV) wherein Z is (C₁-C₁₀)alkyl.

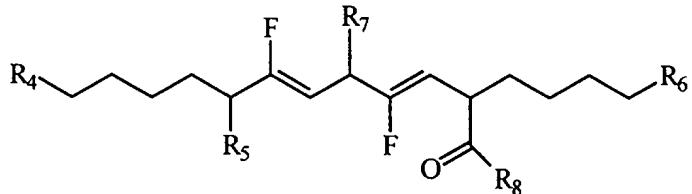
A further preferred compound is a compound of formula (IV) wherein R¹ is indolyl; R² is N(R^a)(R^b); R³ is N(R^c)(R^d); Y is S; Z is hydrogen; and R^a, R^b, R^c, and R^d are each methyl.

25 Yet another preferred compound of formula (IV) includes a compound wherein R¹ is 2-benzimidazolyl; for R² is N(R^a)(R^b); R³ is N(R^c)(R^d); Y is oxo; and Z is N(R^e)(R^f) or a pharmaceutically acceptable salt thereof. Another preferred compound of formula (IV) is a compound wherein R¹ is 2-benzimidazolyl; R² is

N(Me)₂; R³ is N(Me)₂; Y is oxo; and Z is N(Me)₂; or a pharmaceutically acceptable salt thereof.

Another preferred chemokine analog of the invention is a compound of formula (V):

5



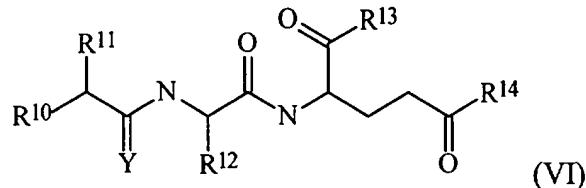
wherein R⁴ is NR_kR_l; wherein R⁵ is NR_mR_n; wherein R⁶ is NR_oR_p; wherein R⁷ is NR_qR_r; wherein R⁸ is hydrogen, hydroxy, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy, NR_sR_t,

10 the amino terminus of an amino acid or the N-terminal residue of a peptide of 2 to about 25 amino acid residues; wherein R_k, R_l, R_o, and R_p are each independently hydrogen, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkanoyl, phenyl, benzyl or phenethyl; wherein R_m are R_n are each independently hydrogen, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₁-C₁₀)alkanoyl, (C₁-C₁₀)alkoxycarbonyl, 9-fluorenylmethoxycarbonyl, phenyl, benzyl, phenethyl, the C-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues; wherein R_q are R_r are each independently hydrogen, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, phenyl, benzyl or phenethyl; wherein R_s are R_t are each independently hydrogen, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, phenyl, benzyl or phenethyl; or a pharmaceutically acceptable salt thereof.

Preferably R_k, R_l, R_o, and R_p are each hydrogen; R_m are R_n are each independently hydrogen, acetyl, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, propoxy, butoxy, *tert*-butoxycarbonyl, 9-fluorenylmethoxycarbonyl, the C-terminal residue of an

amino acid or a peptide of 2 to about 25 amino acid residues; and R_q are R_r are each independently hydrogen, (C₁-C₁₀)alkyl, or (C₃-C₆)cycloalkyl.

A further preferred chemokine analog of the invention is a compound of formula (VI):



5

As used herein, halo is fluoro, chloro, bromo, or iodo. The terms alkyl and alkoxy denote both straight and branched groups, but reference to an individual radical such as "propyl" embraces only the straight chain radical, a branched chain isomer such as "isopropyl" being specifically referred to. Aryl denotes a phenyl

- 10 radical or an ortho-fused bicyclic carbocyclic radical having about nine to ten ring atoms in which at least one ring is aromatic. Heteroaryl encompasses a radical attached via a ring carbon of a monocyclic aromatic ring containing five or six ring atoms consisting of carbon and one to four heteroatoms each selected from the group consisting of non-peroxide oxygen, sulfur, and N(R⁴) wherein R⁴ is absent or
- 15 is hydrogen, (C₁-C₄)alkyl, phenyl or benzyl, as well as a radical of an ortho-fused bicyclic heterocycle of about eight to ten ring atoms derived therefrom, particularly a benz-derivative or one derived by fusing a propylene, trimethylene, or tetramethylene diradical thereto.

It will be appreciated by those skilled in the art that compounds of

- 20 formula (IV), (V), or (VI) including compounds of the invention which are peptides having chiral centers, may exist in and be isolated in optically active and racemic forms. For example, compounds of the invention comprise α -amino acid residues in D or L form, or mixtures thereof. Some compounds may exhibit polymorphism. It is to be understood that the present invention encompasses any racemic, optically-active, polymorphic, or stereoisomeric form, or mixtures thereof, of a compound of
- 25

the invention, which possess the useful properties described herein. It is well known in the art how to prepare optically active forms (for example, by resolution of the racemic form by recrystallization techniques, by synthesis, from optically-active starting materials, by chiral synthesis, or by chromatographic separation using a 5 chiral stationary phase). It is also well known to the art how to determine a compounds ability to inhibit or enhance chemokine-induced activity using the standard tests described herein, or using other tests which are well known in the art.

Specific and preferred values listed herein for radicals, substituents, and ranges, are for illustration only and they do not exclude other defined values or other 10 values within defined ranges for the radicals and substituents. Specifically, (C_1-C_{10}) alkyl can be methyl, ethyl, propyl, isopropyl, butyl, iso-butyl, sec-butyl, pentyl, 3-pentyl, hexyl, heptyl, octyl, nonyl, or decyl; (C_1-C_6) alkyl can be methyl, ethyl, propyl, isopropyl, butyl, iso-butyl, sec-butyl, pentyl, 3-pentyl, or hexyl; (C_1-C_3) alkyl can be methyl, ethyl, or propyl; (C_3-C_6) cycloalkyl can be cyclopropyl, 15 cyclobutyl, cyclopentyl, or cyclohexyl; (C_1-C_{10}) alkoxy can be methoxy, ethoxy, propoxy, isopropoxy, butoxy, iso-butoxy, sec-butoxy, pentoxy, 3-pentoxy, hexyloxy, heptyloxy, octyloxy, nonyloxy, or decyloxy; (C_1-C_6) alkoxy can be methoxy, ethoxy, propoxy, isopropoxy, butoxy, iso-butoxy, sec-butoxy, pentoxy, 3-pentoxy, or hexoxy; (C_1-C_{10}) alkanoyl can be formyl, acetyl, propanoyl, butanoyl, pentanoyl, 20 hexanoyl, heptanoyl, octanoyl, nonanoyl, or decanoyl; (C_1-C_6) alkanoyl can be formyl, acetyl, propanoyl, butanoyl, pentanoyl, or hexanoyl; (C_2-C_6) alkanoyloxy can be acetoxy, propanoyloxy, butanoyloxy, pentanoyloxy, or hexanoyloxy; aryl can be phenyl, indenyl, or naphthyl; and heteroaryl can be furyl, imidazolyl, tetrazolyl, pyridyl, (or its N-oxide), thienyl, benzimidazolyl (or its N-oxide), pyrimidinyl (or its 25 N-oxide), indolyl, or quinolyl (or its N-oxide).

Preferably, the therapeutic agents of the invention are biologically active. The biological activity of a chemokine peptide, peptide variant, analog or derivative thereof, can be measured by methods known to the art, some of which are described hereinbelow. For example, biologically active peptide 3[MCP-1] variants falling

within the scope of the invention have at least about 1%, preferably at least about 10%, more preferably at least about 50%, and even more preferably at least about 90%, the activity of the corresponding native peptide sequence, e.g., peptide 3(1-12)[MCP-1] (SEQ ID NO:1), or the native chemokine, e.g., MCP-1 (SEQ ID NO:16). Thus, a peptide 3 variant, e.g., Leu₄Ile₁₁peptide 3(1-12)[MCP-1], falling within the scope of the invention has an ED₅₀ for inhibition that is at least about 1%, preferably at least about 10%, more preferably at least about 50%, and even more preferably at least about 90%, the maximal activity of peptide 3(1-12)[MCP-1] (SEQ ID NO:1) at 100 μM.

10 Similarly, for example, peptide 2[MCP-1], variants, analogs or derivatives falling within the scope of the invention have an ED₅₀ for a chemokine-like activity which is at least about 1%, preferably at least about 10%, more preferably at least about 50%, and even more preferably at least about 90%, the maximal activity of SEQ ID NO:3 at 100 μM. Alternatively, peptide 2, variants, analogs or derivatives 15 falling within the scope of the invention bind to cells having at least one chemokine receptor with an association constant that is at least about 1%, preferably at least about 10%, more preferably at least about 50%, and even more preferably at least about 90%, the affinity of peptide 2(1-15)[MCP-1] (SEQ ID NO:3) for the same receptor.

20 As used herein, “a chemokine-induced activity” includes, but is not limited to, an activity that is elicited through the binding of a chemokine, a therapeutic agent of the invention or other moiety, e.g., viral protein, to a chemokine receptor, or the binding of a therapeutic agent or other moiety in close physical proximity to the receptor so that the activity is altered. Chemokine receptors include, but are not 25 limited to, CCR1, CCR2a, CCR2b, CCR3, CCR4, CCR5, CCR6, CCR7, CCR8, IL8R1, IL8R2, CC-CKR1, CC-CKR2, CC-CKR3, CXCR1, CXCR2, CXCR3, CX₃CR1 and CXCR4. Chemotide receptors play a role in cell migration, cell activation, viral or parasite entry, release of proinflammatory compounds, and the like.

As used herein, “indications associated with chemokine-induced activity” includes, but is not limited to, atherosclerosis and other forms of local or systemic vasculitis, diseases such as myocardial infarction, stroke and acute ischemia which are secondary to atherosclerosis; hypertension; reperfusion injury (Kumar et al., 5 *Circulation*, 95, 693 (1997)); aortic aneurysms; vein graft hyperplasia; angiogenesis; hypercholesterolemia; congestive heart failure; Kawasaki’s disease; stenosis or restenosis, particularly in patients undergoing angioplasty; pathologically low bone mineral density, such as osteoporosis (Posner et al., *Bone*, 21, 321 (1997)); ulcerative colitis; chronic obstructive pulmonary disease; infection with human 10 immunodeficiency virus (HIV), other lentiviruses or retroviruses with similar mechanisms of cell entry via chemokine receptor(s), or infection with other viruses, e.g., cytomegalovirus (Sozzani et al., *J. Leukoc. Biol.*, 62, 30 (1997)), or viral infection resulting in viral meningitis; organ transplantation, such as acute transplant rejection, allograft rejection and graft versus host disease; transplant vasculopathy; 15 malaria and other consequences of infection by parasites related to plasmodium; asthma; allergic diseases, such as atopy (IgE-mediated components), allergic rhinitis, atopic dermatitis, anaphylaxis, allergic bronchopulmonary aspergillosis (IgE-mediated), and hypersensitivity pneumonitis (high IgG and reactive T cells) (pigeon breeders disease, farmer’s lung disease, humidifier lung disease, malt 20 workers’ lung disease); allergies, including flea allergy dermatitis in mammals such as domestic animals, e.g., dogs and cats, contact allergens including mosquito bites or other insect sting allergies, poison ivy, poison oak, poison sumac, or other skin allergens; urticaria; eczema; pulmonary fibrosis such as idiopathic pulmonary fibrosis; cystic fibrosis; hemolytic uremic syndrome (Van Setten et al., *Pediatr. Res.*, 25 43, 759 (1998)); autoimmune disorders, including, but not limited to, type I diabetes, Crohn’s disease, multiple sclerosis, arthritis, rheumatoid arthritis (Ogata et al., *J. Pathol.*, 182, 106 (1997); Gong et al., *J. Exp. Med.*, 186, 131 (1997)), systemic lupus erythematosus, autoimmune (Hasimoto’s) thyroiditis, autoimmune liver diseases such as hepatitis and primary biliary cirrhosis, hyperthyroidism (Graves’ disease;

thyrotoxicosis), insulin-resistant diabetes, autoimmune adrenal insufficiency (Addison's disease), autoimmune oophoritis, autoimmune orchitis, autoimmune hemolytic anemia, paroxysmal cold hemoglobinuria, Behcet's disease, autoimmune thrombocytopenia, autoimmune neutropenia, pernicious anemia, pure red cell anemia, autoimmune coagulopathies, myasthenia gravis, autoimmune polyneuritis, experimental allergic encephalomyelitis, pemphigus and other bullous diseases, rheumatic carditis, Goodpasture's syndrome, postcardiotomy syndrome, Sjogren's syndrome, polymyositis, dermatomyositis, and scleroderma; eye diseases such as uveitis or blinding Herpes stromal keratitis; liver disease; erhlichiosis or Lyme disease including Lyme arthritis; aberrant hematopoiesis; nephritis due to, for example, autosomal dominant polycystic kidney disease, diabetic nephropathy, IgA nephropathy, interstitial fibrosis, or lupus; as well as other disease states resulting from inappropriate inflammation, either local or systemic, for example, irritable or inflammatory bowel syndrome (Mazzucchelli et al., *J. Pathol.*, 178, 201 (1996)),
10 psoriasis (Gillitzer et al., *Arch. Dermatol. Res.*, 284, 26 (1992); Yu et al., *Lab. Investig.*, 71, 226 (1994)), delayed type hypersensitivity, Alzheimer's disease, chronic pulmonary inflammation, e.g., pulmonary alveolitis and pulmonary granuloma, gingival inflammation or other periodontal disease, and osseous inflammation associated with lesions of endodontic origin (Volejnikova et al., *Am. J. Pathol.*, 150, 1711 (1997)), hypersensitivity lung diseases such as hypersensitivity pneumonitis (Sugiyama et al., *Eur. Respir. J.*, 8, 1084 (1995)), and inflammation related to histamine release from basophils (Dvorak et al., *J. Allergy Clin. Immunol.*, 98, 355 (1996)), such as hay fever, histamine release from mast cells (Galli et al., *Ciba Foundation Symposium*, 147, 53(1989)), or mast cell tumors,
15 types of type 1 hypersensitivity reactions (anaphylaxis, skin allergy, hives, allergic rhinitis, and allergic gastroenteritis); glomerulonephritis (Gesualdo et al., *Kidney International*, 51, 155 (1997)); inflammation associated with peritoneal dialysis (Sach et al., *Nephrol. Dial. Transplant*, 12, 315 (1997)); and pancreatitis.

Other indications falling within the scope of the invention include, but are not limited to, neoplasia, e.g., histiocytoma, glioma, sarcoma, osteosarcoma, osteoma (Zheng et al., J. Cell Biochem., 70, 121 (1998)), melanoma, Kaposi's sarcoma, small cell lung cancer, and ovarian carcinoma as well as myelosuppression and

5 mucositis associated with chemotherapy; brain or spinal cord trauma, such as after disc surgery (Ghirkar et al., J. Neurosci. Res., 46, 727 (1996); Berman et al., J. Immunol., 156, 3017 (1996)); gout; lung disease, e.g., due to respiratory syncytial virus infection of humans, cattle, pigs and the like, or lung injury (Lukacs et al., Adv. Immunol., 62, 257 (1996)); strokes; Loeffler's syndrome; chronic eosinophilic

10 pneumonia; pulmonary fibrosis; wound healing; bacterial infection, e.g., bacterial peritonitis or meningitis; granulomatous diseases such as Mycobacteriosis, Pneumocystosis, Histoplasmosis, Blastomycosis, Coccidiomycosis, Cryptococcosis, Aspergillosis, granulomatous enteritis, Candidiasis, foreign body granulomas and peritonitis, pulmonary granulomatosis, Wegener's granulomatosis (Del Papa et al.,

15 Arthritis Rheum., 39, 758 (1996)), leprosy, syphilis, cat-scratch disease, schistosomiasis (Jacobs et al., Am. J. Pathol., 150, 2033 (1997)), silicosis, sarcoidosis (Iida et al., Thorax, 52, 431 (1997); Car et al., Am. J. Respir. Crit. Care Med., 149, 655 (1994)) and berylliosis; lethal endotoxemia (Zisman et al., J. Clin. Invest., 99, 2832 (1997)); and indications associated with a weak inflammatory

20 response, e.g., which occur in parasitic infection, e.g., *Leishmaniasis* (Moll, Biol. Abs., 104, 21765 (1997)), trypanosome, *Mycobacterium leprae* or *Mycobacterium tuberculosis* infection, helminth infections, such as nematodes (round worms); (Trichuriasis, Enterobiasis, Ascariasis, Hookworm, Strongyloidiasis, Trichinosis, filariasis); trematodes (fluxes) (Schistosomiasis, Clonorchiasis), cestode (tape

25 worms) (Echinococcosis, Taeniasis saginata, Cysticercosis); visceral worms, visceral larva migrans (e.g., *Toxocara*), eosinophilic gastroenteritis (e.g., *Anisaki* spp., *Phocanema* spp.), cutaneous larva migrans (*Ancylostoma braziliense*, *Ancylostoma caninum*), or fungal infection.

In addition, to prevent or treat indications associated with a weak inflammatory response, the agents of the invention, i.e., peptide 2, its variants, analogs and derivatives, may be employed as vaccine adjuvants.

The peptides of the invention may also be useful as contraceptives or to 5 induce abortion, in acute respiratory distress syndrome, and diseases where steroids are routinely used (e.g., relapsing Beheers colitis and asthma).

Also included within the scope of the invention are indications associated with tumor necrosis factor α (TNF α), e.g., rheumatoid arthritis or endotoxemia, or indications associated with elevated levels of TNF α . These indications include, but 10 are not limited to, endotoxic shock; Crohn's disease; fever, and flu-like symptoms; acute interstitial pneumonitis; septic and nonseptic shock; acute respiratory distress syndrome; thromboembolic conditions; bone resorption; arthritis; acute graft versus host disease; cerebral malaria; cachexia of tuberculosis or cancer; lung injury; and idiopathic fibrosis.

15 **I. Identification of Therapeutic Agents Falling within the Scope of the Invention**

Agents useful in the practice of the invention include agents that inhibit or reduce (e.g., chemokine receptor antagonists), or increase, augment or enhance (e.g., chemokine receptor agonists), chemokine-induced activity, e.g., monocyte or macrophage recruitment. These agents can be identified by *in vitro* and *in vivo* 20 assays, such as the assays described hereinbelow. It is recognized that not all agents falling within the scope of the invention can inhibit or enhance chemokine-induced activity *in vitro* and *in vivo*. The therapeutic agents of the invention may be direct receptor binding agonists and/or antagonists, or may act by a different mechanism, e.g., duplex formation of antisense nucleic acid with chemokine mRNA, or by more 25 than one mechanism, so as to result in the alteration of chemokine-induced activity.

A. Peptides, Variants, Derivatives and Analogs

1. *In vitro* chemotaxis

To determine whether an agent inhibits a chemokine-induced activity, such as macrophage recruitment, varying amounts of the agent are mixed with cells in the

presence of a known chemoattractant. For example, a range of known concentrations of an agent, e.g., a chemokine peptide, is incubated with a defined number (e.g., 10^4 - 10^6) of human THP-1 monocyte cells in individual wells of the top compartment of a trans-well plate. Chemokine (such as MCP-1, MIP1 α , IL8 or SDF-1 α), at a concentration known to cause significant migration of THP-1 cells in the trans-well migration assay, is placed in the lower compartment (Figure 1). Cells are then incubated at 37°C for a period sufficient to allow migration, e.g., 4 hours. After incubation, the cells are gently removed from the top of the filter with a pipette, 20 μ l of 20 mM EDTA in simple PBS is added into each top well, and 10 incubated for 20 minutes at 4°C. The filter is carefully flushed with media using a gentle flow, and removed. A standard curve consisting of a two-fold dilution series of THP-1 cells (in 29 μ l) is prepared to accurately quantify the number of cells that have migrated. Migrated cells are stained with 3 μ l of MTT stock dye solution which is added directly into each well (5 mg/ml in RPMI-1640 without phenol red, 15 Sigma Chemical Co.) and incubated at 37°C for 4 hours. The media is carefully aspirated from each well, and the converted dye is solubilized by 20 μ l of DMSO. Absorbance of converted dye is measured at a wavelength of 595 nm using an ELISA plate reader. The number of migrated cells in each well is then determined by interpolation of the standard curve (see also Imai et al., *J. Biol. Chem.*, **272**, 20 15036 (1997)).

Any method suitable for counting cells can be used, for example, counting with a hemocytometer, incubation of the cells with MTT (see above), or FACS analysis. A negative control assay is also performed, using TGF- β or another non-chemokine chemoattractant (e.g., IL1 β or TNF α). To assess whether the agent is 25 cytotoxic, the same concentrations of agent are incubated with THP-1 cells. Agents which 1) are not cytotoxic at levels which inhibit migration, 2) are ineffective at inhibiting the negative control-induced migration, and 3) reduce or inhibit chemokine-induced THP-1 migration, are agents which fall within the scope of the invention.

Agents may also be screened in a chemotactic assay which employs human neutrophils, eosinophils, mast cells, basophils, platelets, lymphocytes or monocytes. For monocytes, 9 mls of fresh blood are transferred to a tube containing 1 ml of 3.8% sodium citrate, and left at room temperature for 15 minutes. Five mls of this 5 anti-coagulated blood are carefully layered over 3.5 ml Polymorphprep® (Nycomed Pharma, Oslo), and centrifuged at 500 g for 35 minutes per the manufacturer's instructions. The top band at the sample/medium interface contains monocytes. The monocytes are carefully removed with a glass pipette, and reconstituted to the original volume (5 ml). The cells are washed with PBS plus 10% fetal calf serum, 10 and centrifuged at 400 g for 10 minutes. The washing step is repeated three times before the cells are counted. Cells are resuspended at 1×10^7 cells/ml in RPMI-1640 + 10% fetal calf serum (FCS). The monocytes are cultured for two days at 37°C in a humidified atmosphere of 5% CO₂.

On day 2, the cells are counted, spun down, and reconstituted to $1 \times 15 10^7$ cells/ml in Gey's balanced salt solution + 1 mg/ml bovine serum albumin (BSA). Chemotaxis is induced in a 48 or 96-well disposable chemotaxis chamber fitted with a 5-8 µm polycarbonate filter for monocytes, neutrophils or eosinophils, or a 3 µm filter for lymphocytes (Uguzzoni et al., *Eur. J. Immunol.*, **25**, 64 (1995); Loetscher et al., *J. Exp. Med.*, **184**, 569 (1996); Weber et al., *J. Immunol.*, 4166 20 (1995)) (PVP free, ChemoTX, Neuroprobe Inc., Cabin John, MD). Twenty-nine µl of chemoattractant or control are added to the lower compartment of each well. The framed filter is aligned with the holes in the corner of the filter frame and placed over the wells. Two and one-half $\times 10^5$ monocytes in 25 µl of Gey's balanced salt solution + 1 mg/ml BSA are added to the upper compartment. The agent is 25 dissolved in Milli Q water and then serially diluted in the Gey's balanced salt solution. In most cases, the serially diluted agent is added to the upper compartment of the chemotaxis chamber. The chamber is incubated at 37°C in a humidified atmosphere of 5% CO₂ for 1.5 hours.

2. Enzyme release

The release of *N*-acetyl- β -D-glucosaminidase from monocytes may be employed to determine whether a therapeutic agent inhibits a cytokine-associated activity. Samples of 1.2×10^6 monocytes in 0.3 ml of prewarmed medium (136 mM NaCl, 4.8 mM KCl, 1.2 mM KH_2PO_4 , 1 mM CaCl_2 , 20 mM Hepes, pH 7.4, 5 mM D-glucose, and 1 mg/ml fatty acid-free BSA) are pretreated for 2 minutes with cytochalasin B (2.7 mg/ml) and then stimulated with a chemokine in the presence or absence of the therapeutic agent. The reaction is stopped after 3 minutes by cooling on ice and centrifugation, and the enzyme activity is determined in the supernatant (Uguzzioni et al., *Eur. J. Immunol.*, **25**, 64 (1995)).

The release of elastase from neutrophils may also be employed to determine whether a therapeutic agent inhibits a cytokine-associated activity (Pereri et al., *J. Exp. Med.*, 1547 (1988); Clark-Lewis et al., *J. Biol. Chem.*, **269**, 16075 (1994)).

3. Cytosolic free Ca^{2+} concentration ($[\text{Ca}^{2+}]_i$) changes

Monocytes, eosinophils, neutrophils and lymphocytes loaded with Fura-2 (0.1 nmol/ 10^5 cells) are stimulated with a chemokine in the presence or absence of the therapeutic agent, and $[\text{Ca}^{2+}]_i$ -related fluorescence changes are recorded (Von Tscharner et al., *Nature*, **324**, 369 (1986)). For example, to determine cytosolic Ca^{2+} concentrations in monocytes, monocytes are incubated with 0.5 μM Fura-2/AM for 30 minutes at 37°C in HEPES-buffered saline (145 mM NaCl, 5 mM KCl, 1 mM MgCl_2 , 10 mM HEPES, and 10 mM glucose), pH 7.4, at 37°C, supplemented with 1% albumin (w/v) and 1 mM CaCl_2 . After loading with Fura-2, the cells are centrifuged for 5 minutes at 300 $\times g$ and then resuspended in buffer containing no added albumin, to a cell density of 1.5×10^6 cells/ml, and kept at room temperature until use. This protocol results in a cytosolic Fura-2 concentration of about 100 μM . Serial dilutions of chemokines in PBS plus 0.1% albumin (w/v) (sterile filtered) are added to aliquots (0.7 ml) of cell suspension. The Fura-2 fluorescence of the monocyte suspension is measured at 37°C in a single excitation, single emission

(500 nm) wavelength Perkin-Elmer LS5 fluorometer. $[Ca^{2+}]_i$ is calculated from changes in fluorescence measured at a single excitation wavelength of 340 nm.

$[Ca^{2+}]_i$ measurements in cells that are stably transformed with a molecularly cloned chemokine receptor which is not expressed in the corresponding non-transformed cells are performed essentially as described above. After loading with Fura-2/AM, cells (1×10^6 /ml) are kept in ice-cold medium (118 mM NaCl, 4.6 mM KCl, 25 mM NaHCO₃, 1 mM KH₂PO₄, 11 mM glucose, 50 mM HEPES, 1 mM MgCl₂, 1 mM CaCl₂, 0.1% gelatin (pH 7.4). Aliquots (2 ml) of cell suspension are prewarmed at 37°C for 5 minutes in 3-ml plastic cuvettes, and fluorescence is measured in a fluorometer (Johnson Foundation Biomedical Group) with magnetic stirring and temperature controlled at 37°C. Excitation is set at 340 nm, and emission is set at 510 nm. $[Ca^{2+}]_i$ is calculated as described above.

For studies in monocytes on cross-desensitization of calcium responses, chemokines are added sequentially with a 2-minute interval, and $[Ca^{2+}]_i$ transients are recorded. The concentrations used in these types of studies vary for each chemokine and are set at levels known to induce the maximal response for $[Ca^{2+}]_i$ mobilization (see Forssmann et al., *FEBS Lett.*, **408**, 211 (1997); Sozzani et al., *J. Leukoc. Biol.*, **57**, 788 (1995); Berkhout et al., *J. Biol. Chem.*, **272**, 16404 (1997)).

4. Chemokine binding and binding displacement

In general, specific binding is calculated as the amount of labeled agent bound in the absence of cold competitor minus the amount of labeled agent bound in the presence of cold competitor. The amount of specific binding in the presence of varied amounts of cold competitor can be used to determine the association constant for the agent, as well as the number of binding sites on the cell for the agent, using, for example, Scatchard Analysis. The agent may be labeled by radiolabeling (e.g., iodination) or with a suitable biochemical tag (e.g., biotin) or by addition of a photoactivatable crosslinking group. Agents with an association constant lower than 100 μ M (i.e., which bind more strongly than an agent with an association constant of 100 μ M) and which have at least about 2,500, preferably at

least about 10,000, and more preferably greater than 25,000, binding sites per cell for at least one cell type which expresses a chemokine receptor, fall under the scope of this invention. THP-1 cells have at least about 5,000 MCP-1 receptors/cell.

For example, monocytes are suspended in RPMI 1640 medium without bicarbonate containing 0.2% bovine serum albumin and 0.1% azide. Radiolabeled chemokine peptide is incubated with 1-2 x 10⁶ cells, e.g., THP-1 cells, in the presence or absence of increasing concentrations of unlabeled chemokine (MCP-1, MCP-3, MCP-4, RANTES or MIP-1 α) for 15 minutes at 37°C in a 96-well plate in a final volume of 0.2 ml (e.g., PBS + 0.5% FCS). After the incubation, 0.5 ml of ice-cold wash buffer (20 mM Tris, 0.5 M NaCl, pH 7.4) is added, and cells are collected onto a polyethyleneimine-treated Whatman GF/C filter using a Brandall cell harvester. Filters are washed with 4 ml of cold wash buffer, and the radioactivity bound to the filters is counted in a γ -counter.

For competition studies, the IC₅₀ is calculated with a curve fitting program (GraFit, Eritacus Software, London), using a four-parameter logistic, $cpm_{bound} = cpm_{max}/(1 + ([L]/IC_{50})^s) + cpm_{ns}$, where cpm_{max} represents the binding without competitor, [L] is the competitor concentration, cpm_{ns} is the non-specific binding, and s is the slope factor. The cpm_{bound} is corrected for “no cell” controls. To obtain the K_d and capacity of binding specific binding, data from homologous displacement experiments are fitted into a single-site ligand binding equation using the GraFit best fit program.

Chemokine binding to cells stably transformed with a molecularly cloned chemokine receptor is performed essentially as described above except that radiolabeled agent is diluted with unlabeled chemokine. Cells are incubated with radiolabeled agent plus or minus unlabeled chemokines for 30 minutes at 37°C (see also, Imai et al., *supra*; Sozzani et al. (1995), *supra*; Berkhout et al., *supra*; WO 97/22698).

5. Binding to the Duffy Antigen Receptor for Chemokines (DARC)

The affinity of the therapeutic agent to DARC may be determined by any method known in the art, e.g., the ability of the agent to inhibit the binding of radio-iodinated MCP-1 to red blood cells (see Example 3). Agents which bind to DARC with a lower association constant (i.e., stronger binding) than they bind to chemokine receptors (i.e., a DARC selectivity ratio of < 1), and which bind to DARC with an association constant lower than 100 μ M, preferably lower than 10 μ M and more preferably lower than 1 μ M, are useful in particular embodiments of the methods of the invention. In contrast, agents which do not bind DARC, or do not bind to DARC with an affinity that is greater than their affinity for chemokine receptors (i.e., a selectivity ratio > 1), are useful in the practice of other embodiments of the methods of the invention.

6. Inhibition of the co-mitogenic activity of chemokines

Many chemokines are co-mitogenic with low concentrations of FCS, e.g., 50 ng/ml MCP-1 + 0.5% FCS is a mitogen for smooth muscle cells. Assays well known to the art for determination of DNA synthesis induced by any known chemokine plus a low concentration (< 5%) of FCS on suitable cells (e.g., smooth muscle cells) in the presence and absence of the agent may be employed to screen agents for such inhibitory activity. See Porreca et al., *J. Vasc. Res.*, 34, 58 (1997), the disclosure of which is incorporated by reference herein.

7. Anti-lentiviral activity

To prepare cell lines that are susceptible to lentiviral infection as a result of the expression of a particular chemokine receptor, a molecularly cloned chemokine receptor is introduced into a cell line that does not otherwise express the chemokine receptor, e.g., HeLa-MAGI (Kimpton and Emerman, *J. Virol.*, 66(5), 3026 (1992)) or U373-MAGI (Harrington and Geballe, *J. Virol.*, 67, 5939 (1993)) cells, by infection with a retroviral vector. Expression of the chemokine receptor on the cell surface is demonstrated by immunostaining live cells using antibody. Expression of the RNA encoding the receptor is demonstrated by RT-PCR analysis. HeLa-MAGI

and U373-MAGI express β -galactosidase after lentiviral infection. Incubation of infected cells with X-gal results in the deposit of a blue stain in these cells.

Infection of the chemokine receptor-stably transformed cell lines with HIV in the presence or absence of agent is performed in 12-well plates with 10-fold serial 5 dilutions of 300 μ l of virus in the presence of 30 μ g/ml DEAE-Dextran as described (Kimpton and Emerman, *supra*). Viral stocks are normalized by ELISA or p24^{gag} (Coulter Immunology) or p27^{gag} (Coulter Immunology) for HIV-1 and HIV-2/SIV, respectively, using standards provided by the manufacturer.

Two days after infection, cells are fixed and stained for β -galactosidase 10 activity with X-gal. The cells are stained for 50-120 minutes at 37°C. The infectious titer is the number of blue cells per well multiplied by the dilution of virus and normalized to 1 ml.

For other methods useful to determine whether an agent inhibits lentiviral 15 infection and/or replication, see also Cocchi et al., *Science*, 270, 1811 (1995), and WO97/22698.

8. Agonists

To determine whether an agent of the invention is a chemokine receptor agonist, varying amounts of a labeled form of the agent, e.g., biotinylated, are mixed with cells that express the receptor, e.g., THP-1 cells express receptors for 20 MCP-1, MIP1 α , SDF-1 α and IL-8, while Jurkat cells express functional receptors for SDF-1. The affinity of the labeled agent for the cells is then determined. Agents that bind to receptors with a reasonable affinity and interact with the receptor by inducing signaling, are within the scope of the invention. While not encompassed by the term “agonist” or “antagonist”, agents that bind to or near the receptor but 25 elicit no response are also within the scope of the invention, and are termed “neutral” agents.

Agents with agonist activity may also be identified using the transwell migration assay, where the cells are placed in the upper compartment (see Figure 1) in the absence of agent, and the agent, e.g. peptide 2[MCP-1], is placed at varying

concentrations in the lower compartment in place of the chemokine. If the agent(s) have agonist activity, more cells are found in the lower compartment at the end of the assay in wells containing the agent(s) than in wells containing inactive control, i.e., agent or medium alone. Preferably, agents having agonist activity also stimulate 5 migration of primary human cells, e.g., monocytes, in a transwell migration assay.

Moreover, weak agonists or neutral agonists (agents which bind to the receptor but do not inhibit binding of native chemokine and its subsequent signaling, nor do they induce signaling themselves) can be identified by screening the agents for ability to displace the binding of HIV gp120, specifically the V3 loop of gp120, 10 to the surface of THP-1 cells or Jurkat cells. Cells are incubated with labeled (for example, radioiodinated) recombinant gp120 protein in an amount effective to bind to the virus receptor, in the presence and absence of various concentrations of the agent(s). Agents which reduce or abolish gp120 binding are agonists or neutral agonists within the scope of the invention.

15 **9. In vivo**

A rapid method to determine whether an agent of the invention inhibits or augments an inflammatory response is to inject a selected chemokine into the skin of an animal in the presence or absence of an agent of the invention. At some later point in time, animals are sacrificed and the number of inflammatory cells in 20 animals exposed to chemokine and the agent is compared to the number of inflammatory cells in animals exposed to chemokine alone, e.g., by quantitative immunofluorescence, relative to control animals.

B. Nucleic acid molecules of the invention

1. Sources of the Nucleic Acid Molecules of the Invention

25 Sources of nucleotide sequences from which the present nucleic acid molecules encoding a chemokine peptide, a variant thereof or the nucleic acid complement thereof, include total or polyA⁺ RNA from any eukaryotic, preferably mammalian, cellular source from which cDNAs can be derived by methods known in the art. Other sources of the DNA molecules of the invention include genomic

libraries derived from any eukaryotic cellular source. Moreover, the present DNA molecules may be prepared *in vitro*, e.g., by synthesizing an oligonucleotide of about 100, preferably about 75, more preferably about 50, and even more preferably about 40, nucleotides in length, or by subcloning a portion of a DNA segment that encodes 5 a particular chemokine.

2. Isolation of a Gene Encoding a Chemokine

A nucleic acid molecule encoding a chemokine can be identified and isolated using standard methods, as described by Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor, NY (1989). For example, reverse- 10 transcriptase PCR (RT-PCR) can be employed to isolate and clone chemokine cDNAs. Oligo-dT can be employed as a primer in a reverse transcriptase reaction to prepare first-strand cDNAs from isolated RNA which contains RNA sequences of interest, e.g., total RNA isolated from human tissue. RNA can be isolated by methods known to the art, e.g., using TRIZOL™ reagent (GIBCO-BRL/Life 15 Technologies, Gaithersburg, MD). Resultant first-strand cDNAs are then amplified in PCR reactions.

"Polymerase chain reaction" or "PCR" refers to a procedure or technique in which amounts of a preselected fragment of nucleic acid, RNA and/or DNA, are amplified as described in U.S. Patent No. 4,683,195. Generally, sequence 20 information from the ends of the region of interest or beyond is employed to design oligonucleotide primers comprising at least 7-8 nucleotides. These primers will be identical or similar in sequence to opposite strands of the template to be amplified. PCR can be used to amplify specific RNA sequences, specific DNA sequences from total genomic DNA, and cDNA transcribed from total cellular RNA, bacteriophage 25 or plasmid sequences, and the like. See generally Mullis et al., Cold Spring Harbor Symp. Quant. Biol., 51, 263 (1987); Erlich, ed., PCR Technology, (Stockton Press, NY, 1989). Thus, PCR-based cloning approaches rely upon conserved sequences deduced from alignments of related gene or polypeptide sequences.

Primers are made to correspond to highly conserved regions of polypeptides or nucleotide sequences which were identified and compared to generate the primers, e.g., by a sequence comparison of other eukaryotic chemokines. One primer is prepared which is predicted to anneal to the antisense strand, and another 5 primer prepared which is predicted to anneal to the sense strand, of a DNA molecule which encodes a chemokine.

The products of each PCR reaction are separated via an agarose gel and all consistently amplified products are gel-purified and cloned directly into a suitable vector, such as a known plasmid vector. The resultant plasmids are subjected to 10 restriction endonuclease and dideoxy sequencing of double-stranded plasmid DNAs.

Another approach to identify, isolate and clone cDNAs which encode a chemokine is to screen a cDNA library. Screening for DNA fragments that encode all or a portion of a cDNA encoding a chemokine can be accomplished by probing the library with a probe which has sequences that are highly conserved between 15 genes believed to be related to the chemokine, e.g., the homolog of a particular chemokine from a different species, or by screening of plaques for binding to antibodies that specifically recognize the chemokine. DNA fragments that bind to a probe having sequences which are related to the chemokine, or which are immunoreactive with antibodies to the chemokine, can be subcloned into a suitable 20 vector and sequenced and/or used as probes to identify other cDNAs encoding all or a portion of the chemokine.

As used herein, the terms "isolated and/or purified" refer to *in vitro* isolation of a DNA or polypeptide molecule from its natural cellular environment, and from association with other components of the cell, such as nucleic acid or polypeptide, 25 so that it can be sequenced, replicated, and/or expressed. For example, "isolated chemokine nucleic acid" is RNA or DNA containing greater than 9, preferably 36, and more preferably 45 or more, sequential nucleotide bases that encode at least a portion of a chemokine, or a variant thereof, or a RNA or DNA complementary thereto, that is complementary or hybridizes, respectively, to RNA or DNA

encoding the chemokine and remains stably bound under stringent conditions, as defined by methods well known in the art, e.g., in Sambrook et al., *supra*. Thus, the RNA or DNA is “isolated” in that it is free from at least one contaminating nucleic acid with which it is normally associated in the natural source of the RNA or DNA

5 and is preferably substantially free of any other mammalian RNA or DNA. The phrase “free from at least one contaminating source nucleic acid with which it is normally associated” includes the case where the nucleic acid is reintroduced into the source or natural cell but is in a different chromosomal location or is otherwise flanked by nucleic acid sequences not normally found in the source cell. An

10 example of isolated chemokine nucleic acid is RNA or DNA that encodes human MCP-1 and shares at least about 80%, preferably at least about 90%, and more preferably at least about 95%, sequence identity with the MCP-1 polypeptide having SEQ ID NO:16.

As used herein, the term “recombinant nucleic acid” or “preselected nucleic acid,” e.g., “recombinant DNA sequence or segment” or “preselected DNA sequence or segment” refers to a nucleic acid, e.g., to DNA, that has been derived or isolated from any appropriate tissue source, that may be subsequently chemically altered *in vitro*, so that its sequence is not naturally occurring, or corresponds to naturally occurring sequences that are not positioned as they would be positioned in a genome

15 which has not been transformed with exogenous DNA. An example of preselected DNA “derived” from a source, would be a DNA sequence that is identified as a useful fragment within a given organism, and which is then chemically synthesized in essentially pure form. An example of such DNA “isolated” from a source would be a useful DNA sequence that is excised or removed from said source by chemical

20 means, e.g., by the use of restriction endonucleases, so that it can be further manipulated, e.g., amplified, for use in the invention, by the methodology of genetic engineering.

Thus, recovery or isolation of a given fragment of DNA from a restriction digest can employ separation of the digest on polyacrylamide or agarose gel by

electrophoresis, identification of the fragment of interest by comparison of its mobility versus that of marker DNA fragments of known molecular weight, removal of the gel section containing the desired fragment, and separation of the gel from DNA. See Lawn et al., Nucleic Acids Res., 9, 6103 (1981), and Goeddel et al., 5 Nucleic Acids Res., 8, 4057 (1980). Therefore, "preselected DNA" includes completely synthetic DNA sequences, semi-synthetic DNA sequences, DNA sequences isolated from biological sources, and DNA sequences derived from RNA, as well as mixtures thereof.

As used herein, the term "derived" with respect to a RNA molecule means 10 that the RNA molecule has complementary sequence identity to a particular DNA molecule.

3. Variants of the Nucleic Acid Molecules of the Invention

Nucleic acid molecules encoding amino acid sequence variants of a chemokine peptide are prepared by a variety of methods known in the art. These 15 methods include, but are not limited to, isolation from a natural source (in the case of naturally occurring amino acid sequence variants) or preparation by oligonucleotide-mediated (or site-directed) mutagenesis, PCR mutagenesis, and cassette mutagenesis of an earlier prepared variant or a non-variant version of the chemokine peptide.

20 Oligonucleotide-mediated mutagenesis is a preferred method for preparing amino acid substitution variants of a chemokine peptide. This technique is well known in the art as described by Adelman et al., DNA, 2, 183 (1983). Briefly, chemokine DNA is altered by hybridizing an oligonucleotide encoding the desired mutation to a DNA template, where the template is the single-stranded form of a 25 plasmid or bacteriophage containing the unaltered or native DNA sequence of the chemokine. After hybridization, a DNA polymerase is used to synthesize an entire second complementary strand of the template that will thus incorporate the oligonucleotide primer, and will code for the selected alteration in the chemokine DNA.

Generally, oligonucleotides of at least 25 nucleotides in length are used. An optimal oligonucleotide will have 12 to 15 nucleotides that are completely complementary to the template on either side of the nucleotide(s) coding for the mutation. This ensures that the oligonucleotide will hybridize properly to the single-stranded DNA template molecule. The oligonucleotides are readily synthesized using techniques known in the art such as that described by Crea et al., Proc. Natl. Acad. Sci. U.S.A., 75, 5765 (1978).

The DNA template can be generated by those vectors that are either derived from bacteriophage M13 vectors (the commercially available M13mp18 and M13mp19 vectors are suitable), or those vectors that contain a single-stranded phage origin of replication as described by Viera et al., Meth. Enzymol., 153, 3 (1987). Thus, the DNA that is to be mutated may be inserted into one of these vectors to generate single-stranded template. Production of the single-stranded template is described in Sections 4.21-4.41 of Sambrook et al., Molecular Cloning: A Laboratory Manual (Cold Spring Harbor Laboratory Press, N.Y. 1989).

Alternatively, single-stranded DNA template may be generated by denaturing double-stranded plasmid (or other) DNA using standard techniques.

For alteration of the native DNA sequence (to generate amino acid sequence variants, for example), the oligonucleotide is hybridized to the single-stranded template under suitable hybridization conditions. A DNA polymerizing enzyme, usually the Klenow fragment of DNA polymerase I, is then added to synthesize the complementary strand of the template using the oligonucleotide as a primer for synthesis. A heteroduplex molecule is thus formed such that one strand of DNA encodes the mutated form of the chemokine, and the other strand (the original template) encodes the native, unaltered sequence of the chemokine. This heteroduplex molecule is then transformed into a suitable host cell, usually a prokaryote such as *E. coli* JM101. After the cells are grown, they are plated onto agarose plates and screened using the oligonucleotide primer radiolabeled with 32-phosphate to identify the bacterial colonies that contain the mutated DNA. The

mutated region is then removed and placed in an appropriate vector for peptide or polypeptide production, generally an expression vector of the type typically employed for transformation of an appropriate host.

The method described immediately above may be modified such that a homoduplex molecule is created wherein both strands of the plasmid contain the mutation(s). The modifications are as follows: The single-stranded oligonucleotide is annealed to the single-stranded template as described above. A mixture of three deoxyribonucleotides, deoxyriboadenosine (dATP), deoxyriboguanosine (dGTP), and deoxyribothymidine (dTTP), is combined with a modified thiodeoxyribocytosine called dCTP-(α S) (which can be obtained from the Amersham Corporation). This mixture is added to the template-oligonucleotide complex. Upon addition of DNA polymerase to this mixture, a strand of DNA identical to the template except for the mutated bases is generated. In addition, this new strand of DNA will contain dCTP-(α S) instead of dCTP, which serves to protect it from restriction endonuclease digestion.

After the template strand of the double-stranded heteroduplex is nicked with an appropriate restriction enzyme, the template strand can be digested with ExoIII nuclease or another appropriate nuclease past the region that contains the site(s) to be mutagenized. The reaction is then stopped to leave a molecule that is only partially single-stranded. A complete double-stranded DNA homoduplex is then formed using DNA polymerase in the presence of all four deoxyribonucleotide triphosphates, ATP, and DNA ligase. This homoduplex molecule can then be transformed into a suitable host cell such as *E. coli* JM101.

For example, a preferred embodiment of the invention is an isolated and purified DNA molecule comprising a preselected DNA segment encoding peptide 3 (1-12)[MCP-1] having SEQ ID NO:1, wherein the DNA segment comprises SEQ ID NO:76, or variants of SEQ ID NO:76, having nucleotide substitutions which are “silent” (see Figure 12). That is, when silent nucleotide substitutions are present in a codon, the same amino acid is encoded by the codon with the nucleotide

substitution as is encoded by the codon without the substitution. For example, valine is encoded by the codon GTT, GTC, GTA and GTG. A variant of SEQ ID NO:79 at the tenth codon in the mature polypeptide (GTC in SEQ ID NO:79) includes the substitution of GTT, GTA or GTG for GTC. Other “silent” nucleotide 5 substitutions in SEQ ID NO:76 which can encode peptide 3 (1-12)[MCP-1] having SEQ ID NO:1 can be ascertained by reference to Figure 12 and page D1 in Appendix D in Sambrook et al., Molecular Cloning: A Laboratory Manual (1989). Nucleotide substitutions can be introduced into DNA segments by methods well known to the art. See, for example, Sambrook et al., *supra*. Likewise, nucleic acid 10 molecules encoding other mammalian, preferably human, chemokines may be modified in a similar manner. Thus, nucleic acid molecules encoding at least a portion of, for example, MCP-2 (SEQ ID NO:80), MCP-3 (SEQ ID NO:81), MCP-4 (SEQ ID NO:100), MIP1 α (SEQ ID NO:82), MIP1 β (SEQ ID NO:83), RANTES (SEQ ID NO:84), SDF1 α (SEQ ID NO:85), IL8 (SEQ ID NO:86), GRO α (SEQ ID 15 NO:87), eotaxin (SEQ ID NO:88), MIG (SEQ ID NO:89), PF-4 (SEQ ID NO:90), I309 (SEQ ID NO:91), HCC-1 (SEQ ID NO:92), C10 (SEQ ID NO:93), CCR-2 (SEQ ID NO:94), ENA-78 (SEQ ID NO:95), GRO β (SEQ ID NO:96), IP10 (SEQ ID NO:97), SDF1 β (SEQ ID NO:98), GRO α (SEQ ID NO:99), MIP3 α , TCA-3, CTAPIII, MARC/FYK, β -thromboglobulin, GCP-2, PBP, HC14, MDC, TECK, 20 PARC, 6Ckine, Fractaline, DC-CK1, LIX, TARC, LARC, MIG, Ck β 8, CCF18/MRP-2, CCIII, CK α 2, H1305, Dvic-1, DGWCC, TCA4, dendrokine, CC2/HCC1, CC3, and MIP1 τ , as well as virally encoded chemokines such as vMIP-I, vMIP-II and vMIP-III, or the complement thereto, may be modified so as to yield nucleic acid molecules of the invention having silent nucleotide substitutions, or to 25 yield nucleic acid molecules having nucleotide substitutions that result in amino acid substitutions (see peptide variants hereinbelow).

C. In vivo studies

To further determine whether a particular agent is useful in the practice of the methods of the invention, an animal model is identified for a human disease.

Transgenic animal models for human disease may also be employed to identify agents useful in the methods of the invention. For example, models of chemokine-induced macrophage recruitment associated with human atherosclerosis include, but are not limited to, mice with a homozygous deletion of the apolipoprotein E (apoE) gene, mice overexpressing human apoB and Watanabe heritable hyperlipidemic rabbits. Models for autoimmune disease include the collagen-induced arthritis in DBA/1 mice and myelin basic protein-induced experimental autoimmune encephalomyelitis. Models for osteoporosis include ovariectomized female rats, mice, monkeys, rats treated with heparin or with glucocorticoids as well as 5 suspension-induced osteoporosis in rats. Models for HIV infection include infection of monkeys with SIV, SIV isolates, HIV or HIV isolates, SCID-Hu mice with HIV or HIV isolates, or rabbits with HIV or HIV isolates. Other animal models for lentiviral infection include cats infected with FIV, horses with EIAV, and goats infected with CAEV (which is also an animal model for arthritis).

10 The efficacy of an agent of the invention may be assessed by measuring the extent of inflammation, or the extent of macrophage infiltration of affected tissues. Macrophage infiltration can be detected by staining tissue sections with antibodies which specifically detect macrophages (e.g., mac-1 antiserum). Inflammation or other symptoms of disease may be detected by measuring appropriate clinical 15 parameters, using techniques which are well known to those skilled in the art. For example, apoE knockout mice are treated with an agent, such as CRD-leu₄ile₁peptide 3, e.g., by intraperitoneal injection, for a period of twelve weeks, while control litter mates receive a suitable control peptide with no known biological activity. At the end of twelve weeks, the animals are sacrificed and the 20 effect of the agent is assessed by measuring the reduction in macrophage recruitment into the vessel wall by quantitative immunohistochemistry using mac-1 antiserum, and by measuring the reduction in the extent of vascular lipid lesion formation by histochemistry using oil red O staining in accordance with Paigen, Arteriosclerosis, 25 10, 316 (1990).

Apo(a) transgenic mice develop lesions when fed a lipid-rich diet. These lesions do not contain any macrophages. In contrast, C57B16 inbred mice develop lipid lesions of similar size and severity to those in apo(a) transgenic mice, but these lesions are rich in infiltrating macrophage. Lesions of apo(a) mice, C57B16 mice, 5 and 6 other strains of mice which develop lipid lesions rich with macrophage, were screened by quantitative immunofluorescence for levels of pro-inflammatory mediators, e.g., TNF- α , MCP-1, MIP-1 α , IL1 β , ICAM-1, VCAM-1, and P-selectin. TNF- α , MIP-1 α , IL1 β , ICAM-1, VCAM-1 and P-selectin were all expressed at identical levels in the apo(a) mouse lesions and the C57B16 lesions. Thus, while 10 these pro-inflammatory mediators may be necessary to infiltration, they are not sufficient alone. In marked contrast, MCP-1 was completely absent from the lesions of apo(a) mice, but expressed at high levels in lesions from all other mouse lines which had macrophage-rich lesions.

Confocal microscopic analysis of sections of blood vessel wall with lesions 15 triple stained with antibodies specific for SM- α -actin (smooth muscle cells; IA4 antibody), macrophages (Mac-1 antibodies) and MCP-1, showed that MCP-1 is not exclusively expressed by macrophage. That is, both smooth muscle cells and macrophages expressed MCP-1. Thus, MCP-1 may be the missing “inflammatory mediator” in the apo(a) mouse model of atherosclerosis. These results suggest that 20 the lack of MCP-1 in apo(a) mice lesions may not be a consequence of the absence of macrophages, but instead contribute to the cause of lack of monocyte infiltration. Moreover, these results provide evidence that the chemokine MCP-1 plays a role in atherosclerotic vascular inflammation. Thus, MCP-1 can provide the basis for analogs which block the recruitment activity of this chemokine.

25 Chemokines other than MCP-1 may also be involved in macrophage recruitment, inflammation and pathogenesis of atherosclerosis, and in other diseases associated with inappropriate proliferation. For example, MIP1 α has been implicated in the inappropriate inflammation in multiple sclerosis. Thus, sequences analogous to peptide 2 and 3 from MIP1 α may be particularly useful to treat or

prevent multiple sclerosis. Therefore, when a particular chemokine is implicated in a particular disease, sequences from that particular chemokine may be especially useful to treat or prevent that disease. Preferred agents falling within the scope of the invention are inhibitors of signaling of more than one chemokine, and preferably of all chemokines. Thus, it may be preferable to prepare chemokine peptide analogs having sequences from a chemokine other than the one(s) associated with a particular disease process. Selection of a particular agent to treat a particular disease may be based on bioavailability, toxicity, DARC binding or other similar criteria.

Other models include, but are not limited to those reported by Lukacs (Adv.

10 Immunol., pp. 257-304, Academic Press (1996)), for lung injury; Lloyd et al. (J.
Leuko. Biol., 185, 1371 (1997)) and Tam et al. (Kid. Int., 49, 715 (1996)), for
nephritis; Volejnikova (Am. J. Pathol., 150, 1711 (1997), for bone; Ghinikar et al.
(J. Neurosci. Res., 46, 727 (1996)) and Ransoholf et al. (J. Leuko. Biol., 62, 645
(1997)), for brain; Kaul et al. (Am. J. Trop. Med. Hyg., 58, 240 (1995)), for malaria;

15 Ajeubar et al. (J. Leuko. Biol., 63, 108 (1998)), for peritonitis; Furukawa et al.
(Lupus, 6, 193 (1997)), for systemic lupus; Suzuki et al. (J. Heart & Lung Transpl.,
16, 1141 (1967)), Abbott et al. (Arch. Surg., 89, 645 (1964)), Corry et al. (Transpl.,
16, 343 (1973)), Dworkin et al. (J. Heart Lung Transpl., 10, 591 (1991)), Laden et
al. (Arch. Path., 93, 240 (1972)) and Mitchell et al. (Transpl., 49, 835 (1990)), for
20 transplants; U.S. Patent No. 5,661,132 for wound healing; Burhardt et al. (Rheum.
Int., 17, 91 (1997)) for autoimmunity; Elson et al. (Gastroenter., 109, 1344 (1998))
for inflammatory bowel disease; Hayes et al. (Arterio. Thromb. Vasc. Biol., 18, 397
(1998)) and Wang et al. (Arterio. Thromb., 11, 1166 (1991)), for cardiovascular
disease; Wegner et al. (Science, 247, 456 (1990) for eosinophilic infiltration into the
25 lung; Brahn (Clinorth and Rel. Res., 265, 42 (1991)), Wooley (Curr. Op. Rheum., 3,
407 (1991)) and Gay et al. (Curr. Op. Rheum., 7, 199 (1995), SCID-human synovial
implant model)) for rheumatoid arthritis); Beamer et al. (Blood, 86, 3220 (1998)),
Nakaguma (Int. J. Exp. Path., 76, 65 (1998)), Nanney et al. (J. Invest. Dermat., 106,
1169 (1996)), Nickoff et al. (AJP, 146, 580 (1995)), Sundberg et al. (Pathobiol., 65,

271 (1997)), and Wolf et al. (Int. J. Dermat., 30, 448 (1998)) for psoriasis; and Conti et al. (Blood, 89, 4120 (1997)), Gonzalo et al. (JCI, 98, 2332 (1996)), Teiyeira et al. (JCI, 100, 1657 (1997)), Ceri et al. (Allergy, 52, 739 (1997)), Freed (Eur. Res. J., 8, 1770 (1998)), Griffiths-Johnson et al. (Meth. Enzy., 288, 241 (1991)), Herz et al. 5 (New Horizons in Allergy Immunoth., 25-32 Plenum Press, 1996) and Kane (Eur. Resp. J., 7, 555 (1991)) for allergy.

II. Preparation of Agents Falling Within the Scope of the Invention

A. Nucleic acid molecules

1. Chimeric Expression Cassettes

10 To prepare expression cassettes for transformation herein, the recombinant or preselected DNA sequence or segment may be circular or linear, double-stranded or single-stranded. A preselected DNA sequence which encodes an RNA sequence that is substantially complementary to a mRNA sequence encoding a chemokine is typically a “sense” DNA sequence cloned into a cassette in the opposite orientation 15 (i.e., 3' to 5' rather than 5' to 3'). Generally, the preselected DNA sequence or segment is in the form of chimeric DNA, such as plasmid DNA, that can also contain coding regions flanked by control sequences which promote the expression of the preselected DNA present in the resultant cell line.

As used herein, “chimeric” means that a vector comprises DNA from at least 20 two different species, or comprises DNA from the same species, which is linked or associated in a manner which does not occur in the “native” or wild type of the species.

Aside from preselected DNA sequences that serve as transcription units for a chemokine, or portions thereof, a portion of the preselected DNA may be 25 untranscribed, serving a regulatory or a structural function. For example, the preselected DNA may itself comprise a promoter that is active in mammalian cells, or may utilize a promoter already present in the genome that is the transformation target. Such promoters include the CMV promoter, as well as the SV40 late promoter and retroviral LTRs (long terminal repeat elements), although many other

promoter elements well known to the art may be employed in the practice of the invention.

Other elements functional in the host cells, such as introns, enhancers, polyadenylation sequences and the like, may also be a part of the preselected DNA.

5 Such elements may or may not be necessary for the function of the DNA, but may provide improved expression of the DNA by affecting transcription, stability of the mRNA, or the like. Such elements may be included in the DNA as desired to obtain the optimal performance of the transforming DNA in the cell.

"Control sequences" is defined to mean DNA sequences necessary for the
10 expression of an operably linked coding sequence in a particular host organism. The control sequences that are suitable for prokaryotic cells, for example, include a promoter, and optionally an operator sequence, and a ribosome binding site. Eukaryotic cells are known to utilize promoters, polyadenylation signals, and enhancers.

15 "Operably linked" is defined to mean that the nucleic acids are placed in a functional relationship with another nucleic acid sequence. For example, DNA for a presequence or secretory leader is operably linked to DNA for a peptide or polypeptide if it is expressed as a preprotein that participates in the secretion of the peptide or polypeptide; a promoter or enhancer is operably linked to a coding
20 sequence if it affects the transcription of the sequence; or a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, "operably linked" means that the DNA sequences being linked are contiguous and, in the case of a secretory leader, contiguous and in reading phase. However, enhancers do not have to be contiguous. Linking is accomplished by
25 ligation at convenient restriction sites. If such sites do not exist, the synthetic oligonucleotide adaptors or linkers are used in accord with conventional practice.

The preselected DNA to be introduced into the cells further will generally contain either a selectable marker gene or a reporter gene or both to facilitate identification and selection of transformed cells from the population of cells sought

to be transformed. Alternatively, the selectable marker may be carried on a separate piece of DNA and used in a co-transformation procedure. Both selectable markers and reporter genes may be flanked with appropriate regulatory sequences to enable expression in the host cells. Useful selectable markers are well known in the art and 5 include, for example, antibiotic and herbicide-resistance genes, such as *neo*, *hpt*, *dhfr*, *bar*, *aroA*, *dapA* and the like. See also, the genes listed on Table 1 of Lundquist et al. (U.S. Patent No. 5,848,956).

Reporter genes are used for identifying potentially transformed cells and for evaluating the functionality of regulatory sequences. Reporter genes which encode 10 for easily assayable proteins are well known in the art. In general, a reporter gene is a gene which is not present in or expressed by the recipient organism or tissue and which encodes a protein whose expression is manifested by some easily detectable property, e.g., enzymatic activity. Preferred genes include the chloramphenicol acetyl transferase gene (*cat*) from Tn9 of *E. coli*, the beta-glucuronidase gene (*gus*) 15 of the *uidA* locus of *E. coli*, and the luciferase gene from firefly *Photinus pyralis*. Expression of the reporter gene is assayed at a suitable time after the DNA has been introduced into the recipient cells.

The general methods for constructing recombinant DNA which can transform target cells are well known to those skilled in the art, and the same 20 compositions and methods of construction may be utilized to produce the DNA useful herein. For example, J. Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press (2d ed., 1989), provides suitable methods of construction.

2. Transformation into Host Cells

25 The recombinant DNA can be readily introduced into the host cells, e.g., mammalian, bacterial, yeast or insect cells by transfection with an expression vector comprising DNA encoding a chemokine or its complement, by any procedure useful for the introduction into a particular cell, e.g., physical or biological methods, to yield a transformed cell having the recombinant DNA stably integrated into its

genome, so that the DNA molecules, sequences, or segments, of the present invention are expressed by the host cell.

Physical methods to introduce a preselected DNA into a host cell include calcium phosphate precipitation, lipofection, particle bombardment, microinjection, 5 electroporation, and the like. Biological methods to introduce the DNA of interest into a host cell include the use of DNA and RNA viral vectors. The main advantage of physical methods is that they are not associated with pathological or oncogenic processes of viruses. However, they are less precise, often resulting in multiple copy insertions, random integration, disruption of foreign and endogenous gene 10 sequences, and unpredictable expression. For mammalian gene therapy, it is desirable to use an efficient means of precisely inserting a single copy gene into the host genome. Viral vectors, and especially retroviral vectors, have become the most widely used method for inserting genes into mammalian, e.g., human cells. Other viral vectors can be derived from poxviruses, herpes simplex virus I, adenoviruses 15 and adeno-associated viruses, and the like.

As used herein, the term "cell line" or "host cell" is intended to refer to well-characterized homogenous, biologically pure populations of cells. These cells may be eukaryotic cells that are neoplastic or which have been "immortalized" *in vitro* by methods known in the art, as well as primary cells, or prokaryotic cells. The cell line 20 or host cell is preferably of mammalian origin, but cell lines or host cells of non-mammalian origin may be employed, including plant, insect, yeast, fungal or bacterial sources. Generally, the preselected DNA sequence is related to a DNA sequence which is resident in the genome of the host cell but is not expressed, or not highly expressed, or, alternatively, overexpressed.

25 "Transfected" or "transformed" is used herein to include any host cell or cell line, the genome of which has been altered or augmented by the presence of at least one preselected DNA sequence, which DNA is also referred to in the art of genetic engineering as "heterologous DNA," "recombinant DNA," "exogenous DNA," "genetically engineered," "non-native," or "foreign DNA," wherein said DNA was

isolated and introduced into the genome of the host cell or cell line by the process of genetic engineering. The host cells of the present invention are typically produced by transfection with a DNA sequence in a plasmid expression vector, a viral expression vector, or as an isolated linear DNA sequence. Preferably, the

5 transfected DNA is a chromosomally integrated recombinant DNA sequence, which comprises a gene encoding the chemokine or its complement, which host cell may or may not express significant levels of autologous or "native" chemokine.

To confirm the presence of the preselected DNA sequence in the host cell, a variety of assays may be performed. Such assays include, for example, "molecular

10 biological" assays well known to those of skill in the art, such as Southern and Northern blotting, RT-PCR and PCR; "biochemical" assays, such as detecting the presence or absence of a particular chemokine, e.g., by immunological means (ELISAs and Western blots) or by assays described hereinabove to identify agents falling within the scope of the invention.

15 To detect and quantitate RNA produced from introduced preselected DNA segments, RT-PCR may be employed. In this application of PCR, it is first necessary to reverse transcribe RNA into DNA, using enzymes such as reverse transcriptase, and then through the use of conventional PCR techniques amplify the DNA. In most instances PCR techniques, while useful, will not demonstrate

20 integrity of the RNA product. Further information about the nature of the RNA product may be obtained by Northern blotting. This technique demonstrates the presence of an RNA species and gives information about the integrity of that RNA. The presence or absence of an RNA species can also be determined using dot or slot blot Northern hybridizations. These techniques are modifications of Northern

25 blotting and only demonstrate the presence or absence of an RNA species.

While Southern blotting and PCR may be used to detect the preselected DNA segment in question, they do not provide information as to whether the preselected DNA segment is being expressed. Expression may be evaluated by specifically identifying the peptide products of the introduced preselected DNA

sequences or evaluating the phenotypic changes brought about by the expression of the introduced preselected DNA segment in the host cell.

B. Peptides, peptide variants, and derivatives thereof

The present isolated, purified chemokine peptides, peptide variants or derivatives thereof, can be synthesized *in vitro*, e.g., by the solid phase peptide synthetic method or by recombinant DNA approaches (see above). The solid phase peptide synthetic method is an established and widely used method, which is described in the following references: Stewart et al., Solid Phase Peptide Synthesis, W. H. Freeman Co., San Francisco (1969); Merrifield, J. Am. Chem. Soc., 85 2149 (1963); Meienhofer in "Hormonal Proteins and Peptides," ed.; C.H. Li, Vol. 2 (Academic Press, 1973), pp. 48-267; Bavaay and Merrifield, "The Peptides," eds. E. Gross and F. Meienhofer, Vol. 2 (Academic Press, 1980) pp. 3-285; and Clark-Lewis et al., Meth. Enzymol., 287, 233 (1997). These peptides can be further purified by fractionation on immunoaffinity or ion-exchange columns; ethanol precipitation; reverse phase HPLC; chromatography on silica or on an anion-exchange resin such as DEAE; chromatofocusing; SDS-PAGE; ammonium sulfate precipitation; gel filtration using, for example, Sephadex G-75; or ligand affinity chromatography.

Once isolated and characterized, derivatives, e.g., chemically derived derivatives, of a given chemokine peptide can be readily prepared. For example, amides of the chemokine peptide or chemokine peptide variants of the present invention may also be prepared by techniques well known in the art for converting a carboxylic acid group or precursor, to an amide. A preferred method for amide formation at the C-terminal carboxyl group is to cleave the peptide from a solid support with an appropriate amine, or to cleave in the presence of an alcohol, yielding an ester, followed by aminolysis with the desired amine.

Salts of carboxyl groups of a peptide or peptide variant of the invention may be prepared in the usual manner by contacting the peptide with one or more equivalents of a desired base such as, for example, a metallic hydroxide base, e.g.,

sodium hydroxide; a metal carbonate or bicarbonate base such as, for example, sodium carbonate or sodium bicarbonate; or an amine base such as, for example, triethylamine, triethanolamine, and the like.

N-acyl derivatives of an amino group of the chemokine peptide or peptide variants may be prepared by utilizing an N-acyl protected amino acid for the final condensation, or by acylating a protected or unprotected peptide. O-acyl derivatives may be prepared, for example, by acylation of a free hydroxy peptide or peptide resin. Either acylation may be carried out using standard acylating reagents such as acyl halides, anhydrides, acyl imidazoles, and the like. Both N- and O-acylation may be carried out together, if desired.

Formyl-methionine, pyroglutamine and trimethyl-alanine may be substituted at the N-terminal residue of the peptide or peptide variant. Other amino-terminal modifications include aminoxyptane modifications (see Simmons et al., Science, 276, 276 (1997)).

In addition, the amino acid sequence of a chemokine peptide can be modified so as to result in a chemokine peptide variant. The modification includes the substitution of at least one amino acid residue in the peptide for another amino acid residue, including substitutions which utilize the D rather than L form, as well as other well known amino acid analogs, e.g., unnatural amino acids such as α , α -disubstituted amino acids, N-alkyl amino acids, lactic acid, and the like. These analogs include phosphoserine, phosphothreonine, phosphotyrosine, hydroxyproline, gamma-carboxyglutamate; hippuric acid, octahydroindole-2-carboxylic acid, statine, 1,2,3,4,-tetrahydroisoquinoline-3-carboxylic acid, penicillamine, ornithine, citruline, α -methyl-alanine, para-benzoyl-phenylalanine, phenylglycine, propargylglycine, sarcosine, ϵ -N,N,N-trimethyllysine, ϵ -N-acetyllysine, N-acetylserine, N-formylmethionine, 3-methylhistidine, 5-hydroxylysine, ω -N-methylarginine, and other similar amino acids and imino acids and tert-butylglycine.

One or more of the residues of the peptide can be altered, so long as the peptide variant is biologically active. For example, for peptide 3[MCP-1] variants,

e.g., Ser₇peptide 3(1-12)[MCP-1], it is preferred that the variant has at least about 10% of the biological activity of the corresponding non-variant peptide, e.g., a peptide having SEQ ID NO:1. Conservative amino acid substitutions are preferred--that is, for example, aspartic-glutamic as acidic amino acids;

5 lysine/arginine/histidine as basic amino acids; leucine/isoleucine, methionine/valine, alanine/valine as hydrophobic amino acids; serine/glycine/alanine/threonine as hydrophilic amino acids. Conservative amino acid substitution also includes groupings based on side chains. For example, a group of amino acids having aliphatic side chains is glycine, alanine, valine, leucine, and isoleucine; a group of

10 amino acids having aliphatic-hydroxyl side chains is serine and threonine; a group of amino acids having amide-containing side chains is asparagine and glutamine; a group of amino acids having aromatic side chains is phenylalanine, tyrosine, and tryptophan; a group of amino acids having basic side chains is lysine, arginine, and histidine; and a group of amino acids having sulfur-containing side chains is

15 cysteine and methionine. For example, it is reasonable to expect that replacement of a leucine with an isoleucine or valine, an aspartate with a glutamate, a threonine with a serine, or a similar replacement of an amino acid with a structurally related amino acid will not have a major effect on the properties of the resulting variant polypeptide. Whether an amino acid change results in a functional peptide can

20 readily be determined by assaying the specific activity of the peptide variant. Assays are described in detail herein.

Conservative substitutions are shown in Figure 13 under the heading of exemplary substitutions. More preferred substitutions are under the heading of preferred substitutions. After the substitutions are introduced, the variants are

25 screened for biological activity.

Amino acid substitutions falling within the scope of the invention, are, in general, accomplished by selecting substitutions that do not differ significantly in their effect on maintaining (a) the structure of the peptide backbone in the area of the substitution, (b) the charge or hydrophobicity of the molecule at the target site,

or (c) the bulk of the side chain. Naturally occurring residues are divided into groups based on common side-chain properties:

- (1) hydrophobic: norleucine, met, ala, val, leu, ile;
- (2) neutral hydrophilic: cys, ser, thr;
- 5 (3) acidic: asp, glu;
- (4) basic: asn, gln, his, lys, arg;
- (5) residues that influence chain orientation: gly, pro; and
- (6) aromatic: trp, tyr, phe.

The invention also envisions peptide variants with non-conservative 10 substitutions. Non-conservative substitutions entail exchanging a member of one of the classes described above for another.

Acid addition salts of the peptide or variant peptide or of amino residues of the peptide or variant peptide may be prepared by contacting the peptide or amine with one or more equivalents of the desired inorganic or organic acid, such as, for 15 example, hydrochloric acid. Esters of carboxyl groups of the peptides may also be prepared by any of the usual methods known in the art.

Moreover, it is also envisioned that the agents of the invention, e.g., chemokine peptides, are modified in a manner that increases their stability *in vivo*, e.g., their half-life or bioavailability. These modified agents are termed 20 "derivatives." Methods to prepare such derivatives are well known to the art. One method to stabilize peptides is to prepare derivatives which are cyclized peptides (see EPA 471,453 (amide bonds), such as that between lysine and aspartic acid side chains; EPA 467,701 (disulfide bonds); EPA 467,699 (thioether bonds). Other modifications which may increase *in vivo* stability are disclosed in Jameson et al. 25 (Nature, 368, 744 (1994)); U.S. Patent No. 4,992,463; U.S. Patent No. 5,596,078 and U.S. Patent No. 5,091,396. A preferred embodiment of the invention is a chemokine peptide or variant that has been cyclized by addition of one or more cysteine residues to the N and/or C terminus of the peptide, as well as peptides which are constructed of the reverse sequence (i.e., reading C-terminal to N-

terminal) of D-form amino acids. A more preferred embodiment of this invention is a peptide which is both cyclized and constructed with the reverse sequence of D-form amino acids, i.e., a CRD derivative.

C. Chemokine analogs

5 Chemokine analogs have properties analogous to those of the corresponding peptide. These analogs can be referred to as "peptide mimetics" or "peptidomimetics" (Fauchere, J. (1986) Adv. Dru Res., 15:29; Veber and Freidinger (1985) TINS p. 392; and Evans et al. (1987) J. Med. Chem., 30:1229, which are incorporated herein by reference) and can be developed with the aid of computerized

10 molecular modeling. These analogs include structures having one or more peptide linkages optionally replaced by a linkage selected from the group consisting of: -CH₂NH-, -CH₂S-, -CH₂-CH₂-, -CH=CH-(cis and trans), -CH=CF-(trans), -CoCH₂-, -CH(OH)CH₂-, and -CH₂SO-, by methods known in the art and further described in the following references: Spatola, A. F. in "Chemistry and

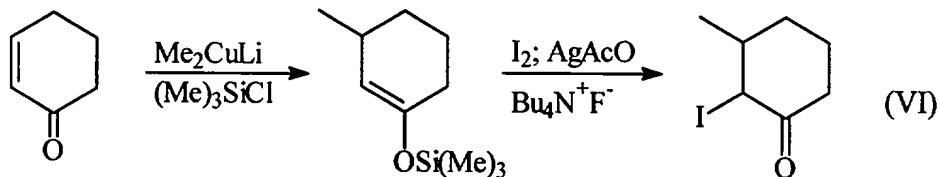
15 Biochemistry of Amino Acids, Peptides, and Proteins," B. Weinstein, eds. Marcel Dekker, New York, P. 267 (1983); Spatola, A. F., Vega Data (March 1983), Vol. 1, Issue 3, "Peptide Backbone Modifications" (general review); Morley, J. S., Trends Pharm. Sci. (1980) pp. 463-468 (general review); Hudson, D. et al., Int. J. Pept. Prot. Res. (1979) 14:177-185 (-CH₂NH-, CH₂CH₂-); Spatola, A. F. et al., Life Sci.

20 (1986) 38:1243-1249 (-CH₂-S); Hann, M. M., J. Chem. Soc. Perkin Trans I (1982) 307-314 (-CH-CH-, cis and trans); Almquist, R. G. et al., J. Med. Chem. (1980) 23:1392-1398 (-COCH₂-); Jennings-White, C. et al., Tetrahedron Lett. (1982) 23:2533 (-COCH₂-); Szelke, M. et al. European Appln. EP 45665 (1982) CA; 97:39405 (1982) (-CH(OH)CH₂-); Holladay, M. W. et al., Tetrahedron Lett. (1983) 24:4401-4404 (-C(OH)CH₂-); and Hruby, V. J., Life Sci. (1982) 31:189-199 (-CH₂S-); each of which is incorporated herein by reference. A particularly preferred non-peptide linkage is -CH₂NH-. Such analogs may have greater chemical stability, enhanced pharmacological properties (half-life, absorption, potency, efficacy, etc.), altered specificity (e.g., a broad-spectrum of biological

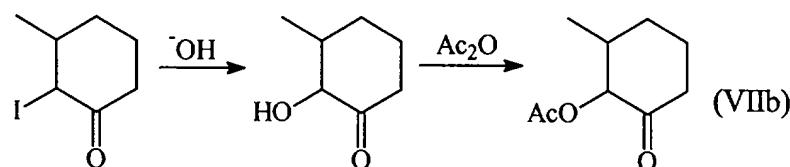
activities), reduced antigenicity, and be economically prepared. Labeling of analogs usually involves covalent attachment of one or more labels, directly or through a spacer (e.g., an amide group), to non-interfering positions(s) on the analog that are predicted by quantitative structure-activity data and/or molecular modeling. Such 5 non-interfering positions generally are positions that do not form direct contacts with the macromolecule(s) to which the analog binds to produce the therapeutic effect. Systematic substitution of one or more amino acids of a consensus sequence with a D-amino acid of the same type (e.g., D-lysine in place of L-lysine) may also be used to generate more stable peptides.

10 1. Isosteres of chemokine tripeptides (a compound of formula (IV))

A compound of formula (IV), wherein Z=CH₃; R=indolyl; Y=O; and X=CH₃, can be prepared from N-tBOC-NinBOC-L-tryptophan-OH and cyclohexenone. For example, 2-cyclohexen-1-one (Aldrich C10,281-4) can be reacted with lithium dimethylcuprate in the presence of trimethylsilyl chloride 15 (Aldrich 38,652-9) (Reaction 1) to trap the enolate intermediate. Lithium dimethylcuprate is prepared from methyl lithium and a copper (I) salt in a 2:1 stoichiometry, prior to use in the reaction, by methods well known to those skilled in the art (e.g., House et al., J. Org. Chem., **40**, 1460 (1975)). The addition of α - β unsaturated ketones by organocuprates is described, for example, in House et al., J. Org. Chem., **31**, 3128 (1966). Similarly, capture of the enolate by trimethyl silyl 20 chloride is described in House et al., J. Org. Chem., **36**, 2361 (1971). The trapped enolate is then resolved to the α -iododerivative, for example, by addition of molecular iodine in the presence of acetoxy-silver and tetrabutylammonium fluoride, according to the method of Rubottom (J. Org. Chem., **44**, 1731 (1979)) to give the 25 trans-disubstituted cyclohexanone of formula (VI),.

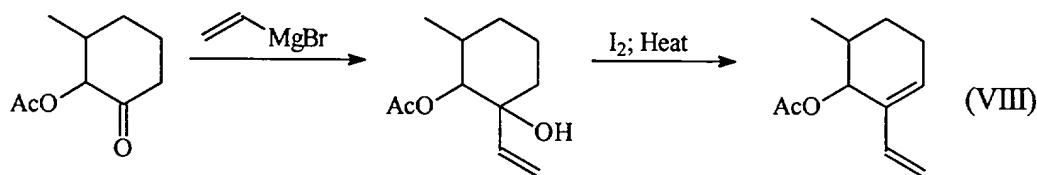


Conversion of the iodide of formula (VI), to a secondary alcohol, and formation of an ester, for example, with acetic anhydride yields a compound of formula (VIIb).

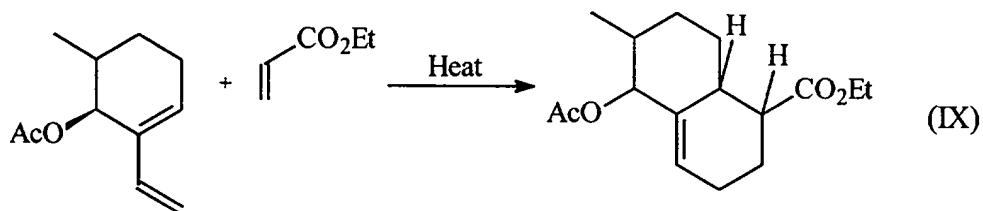


A compound of formula (VIIb) can alternatively be prepared by conversion 5 of the above trimethylsilyl ether enolate to the α -hydroxy ketone followed by formation of the ester, using procedures which are well known in the art.

A compound of formula (VIIb) can be alkylated, for example, with vinyl magnesium bromide under standard conditions, and dehydrated (for example, in the presence of molecular iodine and heat) to yield a diene of formula (VIII):

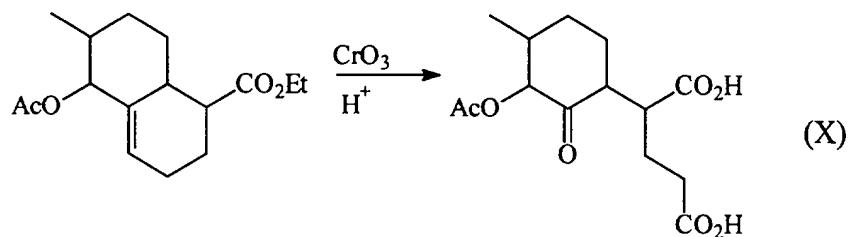


10 Diels-Alder reaction between the diene of formula (VIII) and ethyl acrylate (Aldrich E970-6) gives a stereospecific and regiospecific product of formula IX. For example, the cyclization reaction can be performed by mixing the compound of formula (VIII) and ethyl acrylate in a sealed tube and heating, essentially as described by Green et al. (Adv. Pest Control Res., 3, 129 (1960)).

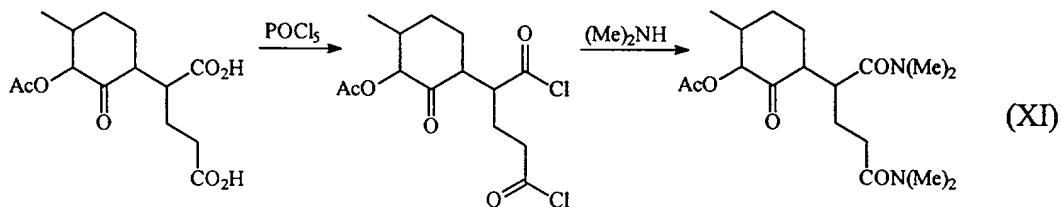


15 Oxidative cleavage of the double bond in a compound of formula (IX) gives a diacid of formula (X). Such an oxidative cleavage may conveniently be carried

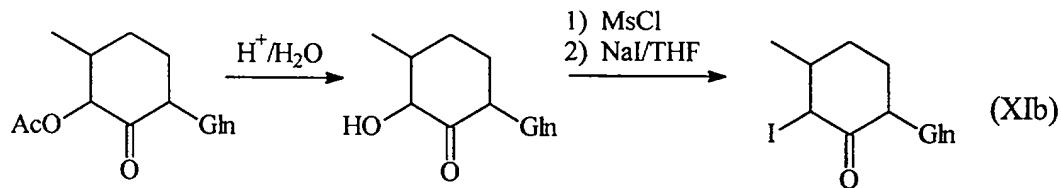
out by ozonolysis or by oxidation with an acid chromate. For example, using CrO_3 in acid, the compound of formula (X) may be prepared, essentially as described by Eschenmoser & Winter, *Science*, 196, 1410 (1977).



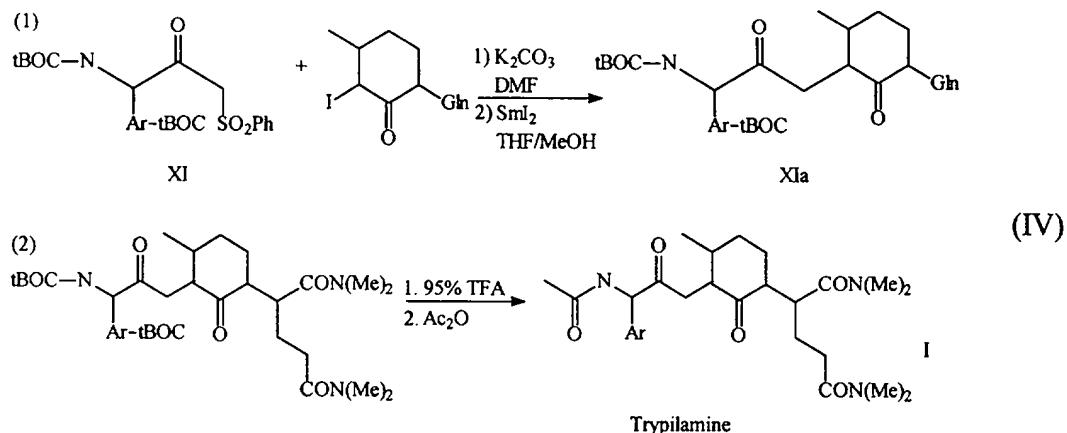
Activation of the diacid with POCl_5 and subsequent reaction with 5 dimethylamine gives a di-amide of formula (XI).



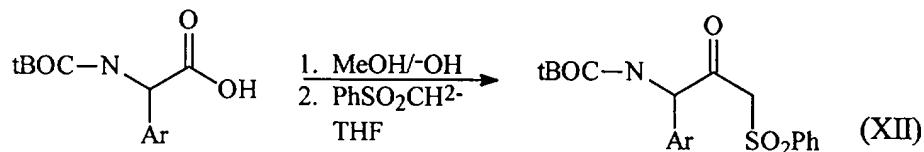
Hydrolysis of the acetoxy group of a compound of formula (XI) followed by formation of the mesylate (or other suitable leaving group) and addition of sodium iodide in THF gives a compound of formula (XIb).



Reaction of a compound of formula (XI) and a compound of formula (XII) in 10 the presence of anhydrous potassium carbonate in dry DMF, essentially as described by Lygo and Rudd (*Tetrahedron Lett.*, 36, 3577 (1995)) followed by removal of the sulfone, for example, using SmI_2 , gives a compound of formula (XII) which can be deprotected and acylated to give a compound of formula (IV) wherein R^2 and R^3 are NMe_2 .



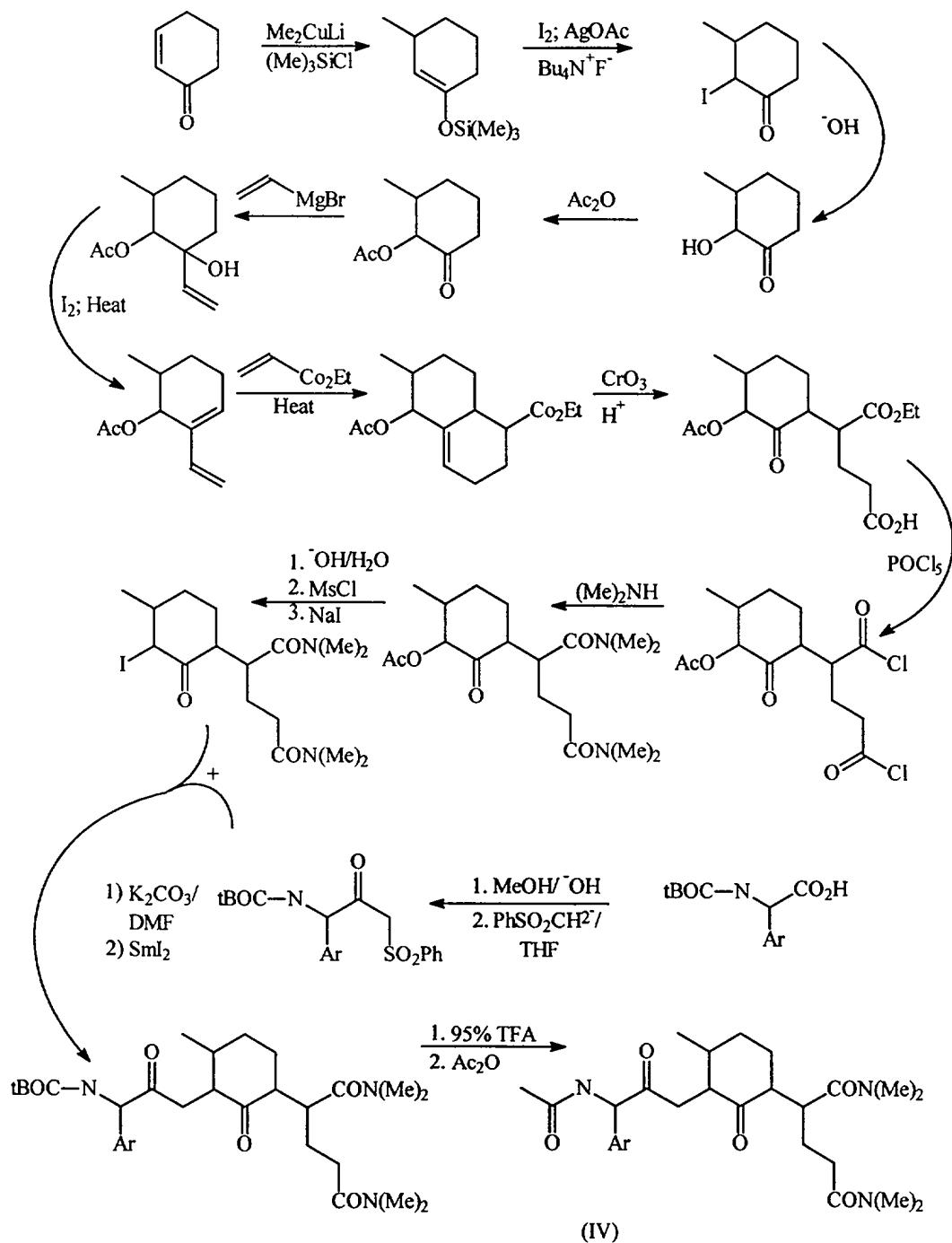
An intermediate of formula (XII) may conveniently be prepared from a protected tryptophan (for example, N- α -tBOC-N_{in}tBOC-L-tryptophan-OH; Novabiochem 04-12-0201) by reaction with the dianion derived of phenylmethylsulfone.



Ar = N-tBOC-Indolyl

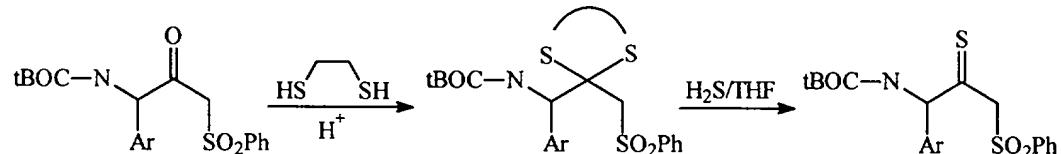
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A preferred synthesis for a compound of formula (IV) is:



Thioketone derivatives ($Y=S$) may be synthesized by insertion of an additional reaction, in which the β -ketosulfone derivative of protected tryptophan is converted to the thioketone derivative. For example, reaction with a dithiol, such as

1,2-ethanedithiol, forms a thioacetal which can be hydrolyzed in the presence of H_2S under anhydrous conditions, to yield the thiol. The conversion may also be carried out using [2,4-bis(4-methoxy-phenyl)-1,3-dithia-2,4-diphosphetane-2,4 disulfide] (Lawesson's Reagent). Reaction of the thiol derivative with the compound of formula (XI) gives a compound of formula (I) wherein Y=S.



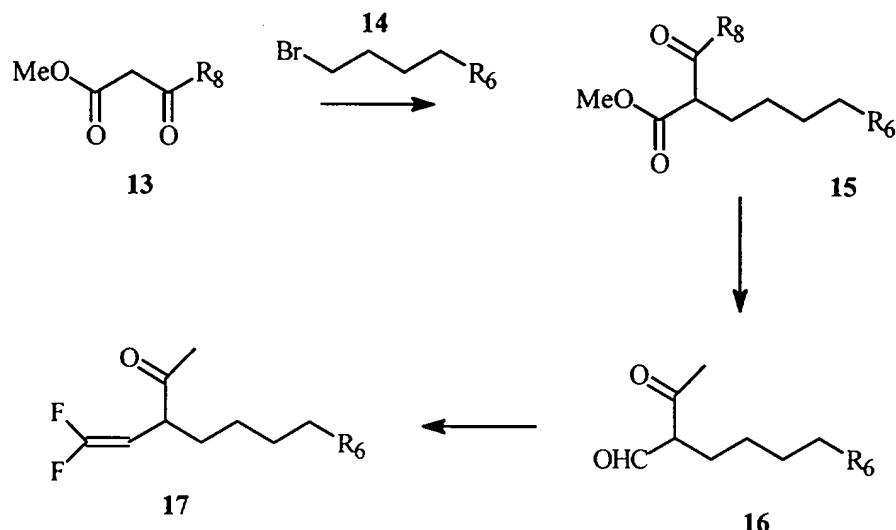
Aryl substituents other than indolyl require preparation of suitably protected β -ketosulfone derivatives of the appropriate amino acid. Where the amino acid is readily available, the reaction can be performed using the appropriate tBOC or Fmoc protected amino acid (phenylalanine and tyrosine, respectively), for example, from Novabiochem. When the amino acid is not readily available (e.g., R=coumaryl), the suitably protected amino acid must first be prepared by methods well established in the art for synthesis of non-standard amino acids (for example, see Yuan and Hruby, *Tetrahedron Lett.*, **38**, 3853 (1997)).

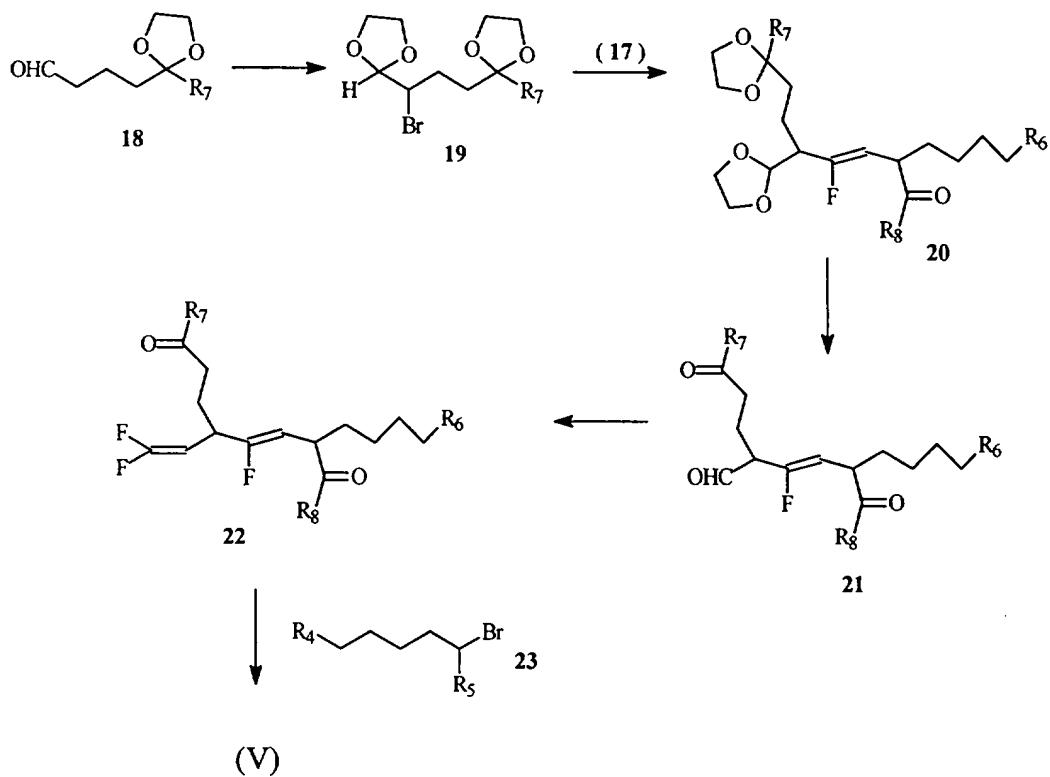
As illustrated below, a compound of formula (V) can conveniently be prepared from an ester of formula 13. Deprotonation with lithium diisopropylamide followed by alkylation with bromide 14 gives a compound of formula 15. Selective reduction of the ester, for example with diisobutylaluminum hydride, gives an aldehyde of formula 16, which can be converted to the difluoroalkene 17 by a Wittig reaction with $\text{PPh}_3=\text{CF}_2$ (Hayashi et al., *Chemistry Letters*, **1980**, pages 935-938).

Aldehyde 18 can be converted to bromide 19 using a procedure similar to that described in Visweswariah et al., *Synthesis*, **1982**, pages 309-310, by treatment with phenyltrimethylammonium tribromide, followed by formation of the acetal under standard conditions. Conversion of the bromide to the corresponding alkylolithium by treatment with n-butyllithium, followed by reaction with difluoride 17, yields a compound of formula 20 (*Chemistry Letters*, **1980**, pages 935-940).

Deprotection under acidic conditions gives aldehyde **21**, which can be reacted with $\text{PPh}_3=\text{CF}_2$ to give trifluoride **22**. Subsequent treatment of **22**, with the alkylolithium derived from bromide **23** yields a compound of formula (V). It will be understood by one skilled in the art that a variety of other known protecting groups can be utilized in the above procedures and that certain protecting groups may be preferred over others depending on the structure of the groups R_4 - R_8 .

5





Other useful chemokine analogs may be identified by the methods described hereinabove. In particular, chemokine analogs that are orally bioavailable, and stable and potent inhibitors of chemokine activity are preferred.

5 D. Targeting of the therapeutic agent

Chemokine peptides, variants, analogs or derivatives thereof may be targeted to a specific therapeutic site by linking the therapeutic agent to a moiety that specifically binds to a cellular component, e.g., antibodies or fragments thereof, lectins, transferrin (for liver targeting) and small molecule drugs, so as to form a therapeutic conjugate. Targeting of the therapeutic agents of the invention can result in increased concentration of the therapeutic agent at a specific anatomic location. Moreover, the linking of a therapeutic agent of the invention to a binding moiety

may increase the stability of the therapeutic agent *in vivo*. For example, an anti-CD4 mimetic that binds to the CD4 receptor may be linked to a therapeutic agent of the invention so as to result in a therapeutic conjugate, a portion of which binds to the HIV co-receptor. This may enhance the ability to target the therapeutic agent to a 5 particular cell type and thus block HIV infection of that cell type.

For neoplasia, anti-tumor antibodies such as NR-LU-10 (anti-carcinoma), NR-ML-5 (anti-melanoma), or anti-CD45 (anti-lymphoma), may be useful to localize the therapeutic agent to a particular type of tumor. For infectious disease, antibodies which recognize a pathogen-specific epitope, such as mAb 17. 41 10 (*Cryptosporidium parvum*), may be employed. To target to joints for treating rheumatoid arthritis, anti-synovium or chondroitin sulfate (e.g., Catalog No. C8035, Sigma Chemical Co., St. Louis, MO) antibodies can be linked to a therapeutic agent of the invention). To treat or prevent asthma or pneumonia, antibodies to the bronchial epithelium may be useful to prepare immunoconjugates for use in the 15 methods of the invention.

Other antibodies useful in targeting a therapeutic agent of the invention to a specific site or cell type include antibodies specific for blood vessels or lymphatics (e.g., Ulex europaeus-I lectin, Catalog No. U4754, Sigma Chemical Co., St. Louis, MO), blood clots or platelets (e.g., Catalog Nos. F9902, F4639, F2506, F8512, 20 Sigma Chemical Co., St. Louis, MO), T cells (e.g., Catalog Nos. C7048 (CD3); C1805 (CD4); C7173 (CD5); and C7298 (CD7), Sigma Chemical Co., St. Louis, MO), brain (e.g., Catalog Nos. S2644 and S2407, Sigma Chemical Co., St. Louis, MO), tumors (e.g., Catalog No. C2331, Sigma Chemical Co., St. Louis, MO), epithelial cells (e.g., Catalog Nos. E6011 and C1041, Sigma Chemical Co., St. 25 Louis, MO), fibroblasts (e.g., Catalog Nos. F4771 and V4630, Sigma Chemical Co., St. Louis, MO), macrophage (e.g., Catalog No. M1919, Sigma Chemical Co., St. Louis, MO), stomach lumen (e.g., Catalog No. M5293, Sigma Chemical Co., St. Louis, MO), neutrophils (e.g., Catalog Nos. N1890 and N1765, Sigma Chemical Co., St. Louis, MO), tendons (e.g., Catalog No. E4013, Sigma Chemical Co., St.

Louis, MO), skin (e.g., Catalog No. K4252, Sigma Chemical Co., St. Louis, MO) mammary tissue or epithelium (e.g., Catalog No. C6930, Sigma Chemical Co., St. Louis, MO) and skeletal muscle (e.g., Catalog Nos. D8281 and D1033, Sigma Chemical Co., St. Louis, MO).

5 To prepare immunoconjugates useful for targeting a malignant or virus-infected cell, an antibody or fragment thereof having a specificity for a surface antigen on a malignant cell or virus-infected is attached to a therapeutic agent of the invention. Preferably, a chemokine peptide or variant thereof is attached via peptide bonds to the carboxy termini regions, e.g., CH3, of antibody heavy chains. The

10 immunoconjugates can be prepared by genetic engineering techniques, i.e., by forming a nucleic acid construct encoding the chimeric immunoconjugate. Preferably, the gene construct encoding the immunoconjugate includes, in 5' to 3' orientation, a DNA segment which encodes a heavy chain variable region, a DNA segment encoding the heavy chain constant region, and a DNA segment coding for

15 the chemokine peptide, peptide variant, or repeats thereof. The fused gene is inserted into an expression vector for transfection of the appropriate recipient cells where it is expressed. The hybrid chain can be combined with a light (or heavy) chain counterpart to form monovalent and divalent immunoconjugates.

The heavy chain constant region for the conjugates can be selected from any

20 of the five isotypes: alpha, delta, epsilon, gamma or mu. Heavy chains or various subclasses (such as the IgG subclasses 1-4) can be used. The light chains can have either a kappa or lambda constant chain. DNA sequences for these immunoglobulin regions are well known in the art (see, e.g., Gillies et al., *J. Immunol. Meth.*, 125, 191 (1989)).

25 In preferred embodiments, the variable region is derived from an antibody specific for the target antigen (an antigen associated with a diseased cell such as a cancer cell or virus-infected cell), and the constant region includes the CH1, CH2 and CH3 domains. The gene encoding the chemokine peptide or variant is joined, e.g., by appropriate linkers, e.g., by DNA encoding (Gly₄-Ser)₃ in frame to the 3' end

of the gene encoding the constant region (e.g., CH3 exon), either directly or through an intergenic region. In certain embodiments, the intergenic region can comprise a nucleotide sequence coding for a proteolytic cleavage site. This site, interposed between the immunoglobulin and the chemokine peptide or variant, can be designed

5 to provide for proteolytic release of the chemokine peptide or variant at the target site. For example, it is well known that plasmin and trypsin cleave after lysine and arginine residues at sites that are accessible to the proteases. Many other site-specific endoproteases and the amino acid sequences they attack are well known.

The nucleic acid construct can include the endogenous promoter and

10 enhancer for the variable region-encoding gene to regulate expression of the chimeric immunoglobulin chain. For example, the variable region encoding genes can be obtained as DNA fragments comprising the leader peptide, the VJ gene (functionally rearranged variable (V) regions with joining (J) segment) for the light chain or VDJ gene for heavy chain, and the endogenous promoter and enhancer for

15 these genes. Alternatively, the gene coding for the variable region can be obtained apart from endogenous regulatory elements and used in an expression vector which provides these elements.

Variable region genes can be obtained by standard DNA cloning procedures from cells that produce the desired antibody. Screening of the genomic library for a

20 specific functionally rearranged variable region can be accomplished with the use of appropriate DNA probes such as DNA segments containing the J region DNA sequence and sequences downstream. Identification and confirmation of correct clones are then achieved by DNA sequencing of the cloned genes and comparison of the sequence to the corresponding sequence of the full length, properly spliced

25 mRNA.

Genes encoding appropriate variable regions can be obtained generally from Ig-producing lymphoid cells. For example, hybridoma cell lines producing Ig specific for tumor associated antigens or viral antigens can be produced by standard somatic cell hybridization techniques. These Ig-producing cell lines provide the

source of variable region genes in functionally rearranged form. The variable region genes are typically of murine origin because the murine system lends itself to the production of a wide variety of Ig's of desired specificity.

The DNA fragment containing the functionally rearranged variable region gene is linked to a DNA fragment containing the gene encoding the desired constant region (or a portion thereof). Ig constant regions (heavy and light chain) can be obtained from antibody-producing cells by standard gene cloning techniques. Genes for the two classes of human light chains and the five classes of human heavy chains have been cloned, and thus, constant regions of human origin are readily available from these clones.

The fused gene encoding the hybrid IgH chain is assembled or inserted into expression vectors for incorporation into a recipient cell. The introduction of gene construct into plasmid vectors can be accomplished by standard gene splicing procedures.

The chimeric IgH chain can be co-expressed in the same cell with a corresponding L chain so that a complete immunoglobulin can be expressed and assembled simultaneously. For this purpose, the heavy and light chain constructs can be placed in the same or separate vectors.

Recipient cell lines are generally lymphoid cells. The preferred recipient cell is a myeloma (or hybridoma). Myelomas can synthesize, assemble, and secrete immunoglobulins encoded by transfected genes and they can glycosylate polypeptide. A particularly preferred recipient cell is the Sp2/0 myeloma which normally does not produce endogenous immunoglobulin. When transfected, the cell will produce only Ig encoded by the transfected gene constructs. Transfected myelomas can be grown in culture or in the peritoneum of mice where secreted immunoconjugate can be recovered from ascites fluid. Other lymphoid cells such as B lymphocytes can be used as recipient cells.

There are several methods for transfecting lymphoid cells with vectors containing the nucleic acid constructs encoding the chimeric Ig chain. A preferred

way of introducing a vector into lymphoid cells is by spheroblast fusion (see Gillies et al., Biotechnol., 7, 798-804 (1989)). Alternative methods include electroporation or calcium phosphate precipitation.

Other useful methods of producing the immunoconjugates include the

5 preparation of an RNA sequence encoding the construct and its translation in an appropriate *in vivo* or *in vitro* system.

Methods for purifying recombinant immunoglobulins are well known. For example, a well known method of purifying antibodies involves protein A purification because of the propensity of protein A to bind the Fc region of

10 antibodies. The antigen binding activity of the purified immunoconjugates can then be measured by methods well known to the art, such as described in Gillies et al. (J. Immunol. Methodol., 125, 191 (1989)). For example, immunoconjugate activity can be determined using antigen-coated plates in either a direct binding or competition assay format.

15 In particular, it is preferred that humanized antibodies are prepared and then assayed for their ability to bind antigen. Methods to determine the ability of the humanized antibodies to bind antigen may be accomplished by any of numerous known methods for assaying antigen-antibody affinity. For example, the murine antibody NR-LU-13 binds an approximately 40 kilodalton glycoprotein expressed

20 on numerous carcinomas. This antigen has been characterized in Varki et al., Cancer Res., 44, 681 (1984); Okabe et al., Cancer Res., 44, 5273 (1989). Thus, it is routine to test the ability of humanized antibodies to bind the NR-LU-13 antigen. Moreover, methods for evaluating the ability of antibodies to bind to epitopes of this antigen are known.

25 Humanized antibodies (or fragments thereof) are useful tools in methods for therapeutic purposes. When determining the criteria for employing humanized antibodies or antibody conjugates for *in vivo* administration for therapeutic purposes, it is desirable that the general attainable targeting ratio is high and that the absolute dose of therapeutic agent delivered to the tumor is sufficient to elicit a

significant tumor response. Methods for utilizing the humanized antibodies can be found, for example, in U.S. Patent Nos. 4,877,868, 5,175,343, 5,213,787, 5,120,526, and 5,202,169.

To target vascular smooth muscle cells (VSMC), VSMC binding proteins, 5 e.g., polypeptides or carbohydrates, proteoglycans and the like, that are associated with the cell membranes of vascular smooth muscle cells can be employed to prepare therapeutic conjugates. In a preferred embodiment, the binding moiety is exemplified by chondroitin sulfate proteoglycans (CSPGs) synthesized by vascular smooth muscle cells and pericytes, and a discrete portion (termed an epitope herein) 10 of the CSPG molecule having an apparent molecular weight of about 250 kD is especially preferred. The 250 kD target is an N-linked glycoprotein that is a component of a larger 400 kD proteoglycan complex. In one presently preferred embodiment of the invention, a vascular smooth muscle binding protein is provided by NR-AN-01 monoclonal antibody (a subculture of NR-ML-05) that binds to an 15 epitope in a vascular smooth muscle CSPG target molecule. The monoclonal antibody designated NR-ML-05 reportedly binds a 250 kD CSPG synthesized by melanoma cells (Morgan et al., U.S. Pat. No. 4,897,255). Smooth muscle cells and pericytes also reportedly synthesize a 250 kD CSPG as well as other CSPGs. NR-ML-05 binding to smooth muscle cells has been disclosed (Fritzberg et al., U.S. 20 Pat. No. 4,879,225). Subculture NR-ML-05 No. 85-41-4I-A2, freeze # A2106, has been deposited with the American Type Culture Collection, Rockville, MD and granted Accession No. HB-9350. NR-ML-05 is the parent of, and structurally and functionally equivalent to, subclone NR-AN-01, disclosed herein. It will be 25 recognized that NR-AN-01 is just one example of a vascular smooth muscle binding protein that specifically associates with the 400 kD CSPG target, and that other binding proteins associating with this target and other epitopes in this target are also useful in the therapeutic conjugates and methods of the invention.

It will be recognized that the inventors also contemplate the utility of human monoclonal antibodies or "humanized" murine antibody as a vascular smooth

muscle binding protein in the therapeutic conjugates of their invention. For example, murine monoclonal antibody may be "chimerized" by genetically recombining the nucleotide sequence encoding the murine Fv region (i.e., containing the antigen binding sites) with the nucleotide sequence encoding a human constant 5 domain region and an Fc region, e.g., in a manner similar to that disclosed in European Patent Application No. 0,411,893 A2. Humanized vascular smooth muscle binding partners will be recognized to have the advantage of decreasing the immunoreactivity of the antibody or polypeptide in the host recipient, which may thereby be useful for increasing the *in vivo* half-life and reducing the possibility of 10 adverse immune reactions. See also, N. Lonberg et al. (U.S. Patent Nos. 5,625,126; 5,545,806; and 5,569,825); and Surani et al. (U.S. Patent No. 5,545,807).

Useful binding peptides for cancer treatment embodiments of the present invention include those associated with cell membrane and cytoplasmic epitopes of cancer cells and the like. These binding peptides localize to the surface membrane 15 of intact cells and internal epitopes of disrupted cells, respectively, and deliver the therapeutic agent for assimilation into the target cells. Minimal peptides, mimetic organic compounds and human or humanized antibodies that localize to the requisite tumor cell types are also useful as binding peptides of the present invention. Such binding peptides may be identified and constructed or isolated in accordance with 20 known techniques. Preferred binding peptides of these embodiments of the present invention bind to a target epitope with an association constant of at least about 10^{-6} M.

Methods useful to prepare antibody-peptide conjugates are well known to the art. See, for example U.S. Patent No. 5,650,150, the disclosure of which is 25 incorporated by reference herein. Representative "coupling" methods for linking the therapeutic agent through covalent or non-covalent bonds to the targeting moiety include chemical cross-linkers and heterobifunctional cross-linking compounds (i.e., "linkers") that react to form a bond between reactive groups (such as hydroxyl, amino, amido, or sulphydryl groups) in a therapeutic agent and other reactive groups

(of a similar nature) in the targeting moiety. This bond may be, for example, a peptide bond, disulfide bond, thioester bond, amide bond, thioether bond, and the like. In one illustrative example, conjugates of monoclonal antibodies with drugs have been summarized by Morgan and Foon (Monoclonal Antibody Therapy to

5 Cancer: Preclinical Models and Investigations, Basic and Clinical Tumor Immunology, Vol. 2, Kluwer Academic Publishers, Hingham, MA) and by Uhr, J. of Immunol. 133:i-vii, 1984). In another illustrative example where the conjugate contains a radionuclide cytostatic agent, U.S. Patent No. 4,897,255, Fritzberg et al., incorporated herein by reference, is instructive of coupling methods that may be

10 useful. In one embodiment, the therapeutic conjugate contains a vascular smooth muscle binding protein coupled covalently to a chemokine peptide or variant. In this case, the covalent bond of the linkage may be formed between one or more amino, sulphydryl, or carboxyl groups of the vascular smooth muscle binding protein and the chemokine peptide or variant.

15 In a preferred embodiment of the invention, an antibody conjugate is used in pretargeting methods. Essentially, such pretargeting methods are characterized by an improved targeting ratio or increased absolute dose to the target cell sites in comparison to conventional cancer diagnosis or therapy. A general description of pretargeting methods may be found in U.S. Patent No. 4,863,713, 5,578,287, and

20 5,630,996. Typical pretargeting approaches are summarized below.

Pretargeting methods are of two general types: three-step pretargeting methods and two-step pretargeting methods. A three-step pretargeting protocol includes the administration of a targeting moiety-ligand conjugate, which is allowed to localize at a target site and to dilute in the circulation. This is followed by

25 administration of an anti-ligand which binds to the targeting moiety-ligand conjugate and clears unbound targeting moiety-ligand conjugate from the blood, as well as binds to targeting moiety-ligand conjugate at the target site. Thus, the anti-ligand fulfills a dual function by clearing targeting moiety-ligand conjugate not bound to the target site as well as attaches to the target site to form a targeting

moiety-ligand : anti-ligand complex. Finally, a therapeutic agent-ligand conjugate that exhibits rapid whole body clearance is administered.

When the therapeutic agent-ligand conjugate in circulation comes into close proximity to the targeting moiety-ligand : anti-ligand complex bound to the target site, the anti-ligand portion of the complex binds to the ligand portion of the circulating therapeutic agent-ligand conjugate, thus producing a targeting moiety-ligand : anti-ligand : ligand-therapeutic agent "sandwich" at the target site. Furthermore, because the unbound therapeutic agent is attached to a rapidly clearing ligand (rather than a slowly clearing targeting moiety, such as antibody or antibody fragment), this technique provides decreased non-target exposure to the active agent.

Alternatively, two-step pretargeting methods eliminate the step of administering the above identified anti-ligand. These "two-step" procedures feature targeting moiety-ligand or targeting moiety-anti-ligand administration, followed by the administration of a therapeutic agent which is conjugated to the opposite 15 member of the ligand/anti-ligand pair.

As an optional step in the two-step pretargeting method, ligand or anti-ligand, designed specifically to provide a clearance function, is administered to facilitate the clearance of circulating targeting moiety-ligand or targeting moiety-anti-ligand. Thus, in the two-step pretargeting approach, the clearing agent does not 20 become bound to the target cell population, either directly or through the previously administered target cell bound targeting moiety-anti-ligand or targeting moiety-ligand conjugate.

A targeting moiety in a pretargeting method binds to a defined target cell population, such as tumor cells. Preferred targeting moieties useful in this regard 25 are antibodies (polyclonal or monoclonal), such as human monoclonal antibodies, or "humanized" murine or chimeric antibodies. Some examples of humanized antibodies include those that are CHO produced, produced in hosts such as plant (for example corn, soybean, tobacco, and the like), insect, mammalian, yeast, and bacterial. The humanized antibodies may be those that bind to the antigen bound by

antibody NR-LU-13. Preferably, the humanized antibody may not possess N-linked glycosylation or its N-linked glycosylation has been modified post expression to reduce immunogenicity or toxicity.

Ligand/anti-ligand pairs suitable for use in targeting protocols include

- 5 biotin/avidin or streptavidin, haptens and epitopes/antibody, fragments or analogs thereof, including mimetics, lectins/carbohydrates, zinc finger proteins/dsDNA fragments, enzyme inhibitors/enzymes; and analogs and derivatives thereof.
- Preferred ligands and anti-ligands bind to each other with an affinity of at least about $K_A \geq 10^9 M^{-1}$ or $K_D \leq 10^{-9} M$. Biotin/avidin or streptavidin is a preferred ligand/anti-
- 10 ligand pair.

In general, such pretargeting methods preferably include the administration of an anti-ligand that provides a clearance function. The clearance is probably attributable to cross-linking and/or aggregation of conjugates that are circulating in the blood, which leads to complex/aggregate clearance by the recipient's RES (reticuloendothelial system). The anti-ligand clearance of this type is preferably accomplished with a multivalent molecule. However, a univalent molecule of sufficient size to be cleared by the RES on its own could also be employed.

Alternatively, receptor-based clearance mechanisms, e.g., Ashwell receptor or other receptors, may be exploited by addition of hexose residues, such as galactose or mannose residues, to provide for clearance of the anti-ligand, anti-ligand conjugate or humanized antibody via the liver. Such clearance mechanisms are less dependent upon the valency of the clearing agent than the RES complex/aggregate clearance mechanisms described above.

For example, if the targeting moiety-ligand or targeting moiety-anti-ligand has been derivatized to provide for clearance (i.e., addition of a hexose residue) a clearing agent should not be necessary. Preferred clearing agents are disclosed in U.S. Patent Nos. 5,624,896 and 5,616,690; as well as PCT Application Publication Number WO 95/15978.

One skilled in the art, based on the teachings herein and the applications referenced herein, can readily determine an effective therapeutic effective dosage and treatment protocol. This will depend upon factors such as the particular selected therapeutic agent, route of delivery, the type of target site(s), affinity of the targeting moiety for target site of interest, any cross-reactivity of the targeting moiety with normal tissue, condition of the patient, whether the treatment is effected alone or in combination with other treatments, among other factors.

For example, in the case of humanized antibody - avidin or streptavidin conjugates in pretargeting strategies, a suitable dosage ranges from about 10 to 10 2500 mg, more preferably from about 50 to 1500 mg, and most preferably from about 100 to 800 mg. The dosage of the ligand-therapeutic agent conjugate, generally ranges from about 0.001 to about 10 mg and more preferably from about 0.1 to 2 mg.

In general, such pretargeting methods include the administration of a 15 clearing agent. The dosage of the clearing agent is an amount which is sufficient to substantially clear the previously administered conjugate from the circulation, i.e., at least about 50%, more preferably at least about 90%, and most preferably approaching or at 100%. In general, the clearing agent is administered several days after administration of the humanized antibody - streptavidin conjugate, preferably 20 about 1 to 5 days after, more preferably at least about 1 to 2 days after. Generally, the determination of when to administer the clearing agent depends on the target uptake and endogenous clearance of targeting moiety conjugate. Particularly preferred clearing agents are those which provide for Ashwell receptor mediated clearance, such as galactosylated proteins, e.g., galactosylated biotinylated human 25 serum albumin (HSA) and small molecule clearing agents containing galactose and biotin. In the case of HSA based clearing agents, a typical dosage of the clearing agent will range from about 100 to 1000 mg, and more preferably about 200-500 mg. If a clearing agent is administered, the ligand-therapeutic agent conjugate is preferably administered about 2 to 12 hours after.

The conjugates may be administered by known methods of administration.

Known methods of administration include, by way of example, intraperitoneal injection, intravenous injection, intramuscular injection, intranasal administration, among others. Intravenous administration is generally preferred.

5 III. Indications Amenable to Treatment by the Agents of the Invention

The agents of the invention are useful to treat a mammal afflicted with, to inhibit in a mammal at risk of, or to augment in a mammal at risk of, an indication associated with chemokine-induced activity, such as aberrant or pathological inflammatory processes. The chemokines participate in a broad range of

10 inflammatory processes, both physiological and pathological. Thus, broad specificity chemokine inhibitors may be useful to treat or prevent a wide range of inflammatory diseases. Moreover, the use of rationally designed chemokine inhibitors, i.e., inhibitors with relative specificity for various chemokines, may reduce or inhibit side-effects associated with chronic therapies of broad spectrum

15 chemokine inhibitors. Thus, these inhibitors may be designed to treat particular diseases, thereby minimizing side effects resulting from disrupting unrelated physiological processes.

Atherosclerosis. Development of atherosclerosis is a complex process involving smooth muscle cells, endothelial cells and inflammatory cells, and, in particular, monocyte-derived tissue macrophages, B or T cells. Once endothelial cells are activated, they express adhesion molecules important for the extravasation of inflammatory cells. For example, in the TGF β 1 knockout (-/-) mouse, the absence of this cytokine resulted in endothelial cell activation. The activated endothelial cells express, among other adhesion molecules, E-selectin, P-selectin, and ICAM-1, which in turn participate in the extravasation of leukocytes. Potent pro-inflammatory cytokines were also expressed at the sites of incipient vascular lesions. TNF- α , IL-1, as well as several chemokines including IL-8 and MCP-1, have been detected at elevated levels in atherosclerotic lesions. Results described

hereinabove show that the chemokine MCP-1 in particular plays a role in atherosclerotic vascular inflammation.

It is now well accepted that the acute stability of vascular lesions is a more important determinant of short-term, e.g., less than several years, risk of myocardial infarction than is total plaque burden. The degree of macrophage infiltration is probably the major determinant of relative plaque stability. At least two factors contribute to plaque stability: macrophages secrete an excess of matrix-degrading enzymes (such as the matrix metalloproteinases) over their inhibitors, resulting in the loss of extracellular matrix (ECM) in the macrophage-rich shoulder and fibrous cap regions, a common feature of unstable or ruptured plaques; and macrophage-derived foam cells become necrotic, possibly in response to toxic oxidative metabolites of lipids, resulting in a lipid-filled extracellular pool which further destabilizes the local vessel wall architecture.

Inhibitors of chemokine action, and in particular inhibitors of MCP-1, may improve plaque stability and thus rapidly reduce the risk of myocardial infarction, without necessarily reducing the total atherosclerotic plaque burden. In particular, the agents of the invention may decrease lipid lesion formation and/or lipid lesion progression as well as increasing plaque stability (Boring et al., *Nature*, 394, 894 (1998)). Thus, agents of the invention, e.g., peptide 3(1-12)[MCP-1] (SEQ ID NO:1), KQK, peptide 3[7-12] (SEQ ID NO:9), as well as variants, e.g., Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14), or derivatives thereof, may be useful to treat and/or prevent unstable angina pectoris, atherosclerosis, as well as other diseases characterized by local or systemic vasculitis, as well as the symptoms and diseases which occur secondarily to the vessel wall inflammation such as myocardial infarction.

Moreover, the agents of the invention are also useful in combination with lipid lowering agents, such as the statins, or TGF-beta elevating agents (see, for example, WO 96/40098, the disclosure of which is incorporated by reference herein).

Osteoporosis. Low bone mineral density, often categorized as osteoporosis, results from an imbalance between bone matrix deposition by osteoblasts and its subsequent resorption by osteoclasts. The balance between these two dynamic processes determines bone density. One strategy to increase bone density has been 5 the use of analogs of tamoxifen, such as raloxifene, which mimic the effects of estrogen on bone and thus, promote osteoblast differentiation (increasing bone matrix deposition) and inhibit osteoclast recruitment (decreasing resorption). An alternative strategy is to decrease matrix resorption by directly inhibiting the mechanism by which osteoclasts are recruited to the bone. Measurement of bone 10 matrix degradation products (such as the N-terminal and C-terminal telopeptides of collagen as well as pyridinium cross-links) in plasma and urine confirm that bone resorption is increased in osteoporosis, and hence inhibition of osteoclast activity is likely to prove an effective therapeutic strategy.

Unlike osteoblasts, which are locally derived, osteoclasts are continuously 15 recruited to bone as precursor cells which circulate in the monocyte fraction, and which may be identical to monocytes. Once recruited, the precursors differentiate into osteoclasts which then resorb matrix until they die by apoptosis. Thus, the number of osteoclasts in bone tissue (and hence the osteoclast activity) can be rapidly regulated by modulating the osteoclast recruitment process.

20 A number of lines of evidence now suggest that the monocyte recruitment into bone is a molecular parallel of the pathological monocyte recruitment into the blood vessel wall that occurs during atherogenesis. In particular, the chemokine MCP-1 is implicated in both processes. Thus, MCP-1 inhibitors may act to reduce monocyte recruitment and thus decrease osteoclast recruitment and/or decrease the 25 number of cells differentiating into osteoclasts, which would result in a rapid increase in bone density, for example, over a period of weeks rather than years. The ability of the present therapeutic agents to increase bone density contrasts with existing drugs which prevent a further decrease in bone density but do not increase bone density. Therefore, peptide 3, e.g., peptide 3(7-12)[MCP-1], and variants (e.g.,

Leu₄Ile₁₁peptide 3(1-12)[MCP-1]) and derivatives (e.g., CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1]) thereof, may be useful to inhibit or prevent low bone density. In particular, derivatives with specificity for CC chemokines, such as CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1], are preferred agents for the treatment of 5 osteoporosis.

HIV Infection and AIDS. In addition to the CD4 receptor, additional cell surface molecules (termed co-receptors) are required for the productive infection of a cell by HIV isolates. HIV isolates can be divided into two subtypes, which depend on whether they can infect monocyte/macrophages (M-tropic strains) or helper T 10 lymphocytes (T-tropic strains). Experiments with chemokine ligands suggest that the chemokine receptors function as the HIV co-receptors: MIP1 α and RANTES inhibited the infection of monocytes with M-tropic strains (but not infection of T-cells by T-tropic strains), while SDF-1 inhibited T cell infection (but not monocyte infection). Further molecular analyses confirmed that the MIP1 α /RANTES receptor 15 CCR-5 is the HIV co-receptor on monocytes while the SDF-1 receptor CXCR-4 (also termed LESTR and fusin) is the co-receptor on T-cells. Early in infection, M-tropic virus predominates, a virus which is non-syncytium forming, less virulent and does not deplete T-cells. At a later time, selection favors conversion to the more virulent, syncytium forming T-tropic strain, a strain which depletes helper T cells 20 and leads to acquired immunodeficiency (AIDS). It is possible that the virions can use other chemokine receptors, although at lower efficiency. Thus, to provide an effective agent to inhibit HIV, the agent preferably inhibits virus binding to more than one receptor, i.e., an agent would have to have broad specificity for chemokine receptors.

25 Genetic studies have identified a mutation in CCR5 which renders individuals essentially immune to HIV infection. This mutation, termed CCR5 Δ 32, results in a truncated mRNA for CCR-5. The expression of the truncated CCR-5 does not produce any detectable CCR-5 protein on the cell surface. Individuals homozygous for this deficiency have been reported to be entirely resistant to HIV

infection, even under exposure to extremely high viral challenge, although there is now a single report of a homozygous mutant individual seropositive for HIV infection. Thus, these observations demonstrate that effective blockade of the CCR-5 receptor may effectively prevent infection. Moreover, CCR-5 mediated 5 chemokine signaling does not have a crucial role in normal physiology, since CCR-5 Δ 32 homozygotes have no detectable phenotype other than HIV resistance.

Therefore, inhibitors of chemokine receptors, such as peptide 3, its variants, 10 analogs or derivatives, may inhibit HIV infection as these agents have broad specificity. As described hereinbelow (Example 5), peptide 3[MCP-1] inhibited HIV binding and infection of Jurkat cells and macrophage. A preferred agent to prevent or inhibit HIV infection and/or replication is CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1]. In particular, peptide 3, its variants, analogs or derivatives, e.g., CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1], may be especially useful to inhibit 15 infection of M-tropic strains of HIV.

Peptide 2, its variants, analogs or derivatives, are also useful to prevent or 20 inhibit HIV infection and/or replication, as peptide 2 inhibited HIV replication in T cells and macrophage. Preferred therapeutic agents have decreased Duffy binding and increased co-receptor affinity (in at least about the nM range) (see Example 5) relative to the corresponding chemokine or peptide having the native or wild-type sequence. Preferably, Peptide 2, its variants, analogs or derivatives, e.g., LRD derivatives, are useful to inhibit T-tropic strains of HIV.

Thus, a combination of peptide 3, its variants, analogs or derivatives, and peptide 2, its variants, analogs or derivatives, may be particularly useful to prevent or treat HIV infection.

25 Thus, these agents are useful for the treatment, as well as the prevention, of both HIV seropositives and of progression of seropositive patients to AIDS, when used, either alone, in combination, or in combination with other anti-viral therapies. When used in combination, it is preferred that an infected individual is pre-treated with viral inhibitors (such as a cocktail of reverse transcriptase and viral protease

inhibitors) and then given doses of a general chemokine inhibitor, preferably peptide 3, peptide 2, their variants or derivatives, more preferably peptide 2[MIP1 α], its analogs or derivatives. Moreover, since resistance to other therapies (such as protease inhibitors or reverse transcriptase inhibitors) arise because of viral

5 replication, agents which reduce virus infectivity may drastically increase the success of these existing therapies. Specifically, unlike all currently exploited therapeutic targets such as reverse transcriptase or the viral protease, chemokine agonists and/or antagonists target the susceptible cell rather than the virus itself. Although the virus can rapidly mutate to generate strains resistant to the virus-

10 targeted agents, cells mutate less readily and are under less or no selective pressure to mutate. The extent to which the mutations in the HIV virus must occur to circumvent the use of a chemokine co-receptor is likely to be much greater than the mutations necessary to render a reverse transcriptase resistant to a reverse transcriptase inhibitor. Thus, the administration of chemokine analogs is likely to

15 prove effective either alone or in combination with the virus-targeted therapies. Furthermore, chemokine inhibitors may have limited side effects *in vivo*, i.e., limited physiological impact, and therefore have a good therapeutic index when used *in vivo*.

Malaria. Malaria is caused by intracellular parasites of the plasmodium group. The main cellular target of the parasite is the red blood cell, and the mechanism of infection involves interaction of a plasmodium surface protein and the Duffy Antigen Receptor for Chemokines (DARC) for at least one common plasmodium species, *P. vivax*. Studies have indicated that normal surface presentation of DARC on the red blood cell surface is necessary for plasmodium entry. Consequently, inhibitors of DARC function may be useful anti-malarial agents. Hence, peptide 3, or preferably a derivative showing higher affinity DARC binding such as Ser₇Glu₈Glu₉peptide 3(1-12)[MCP-1], or more preferably, a peptide which shows high affinity DARC binding and does not inhibit chemokine activity

such as peptide 2[MCP-1], peptide 2[MGSA], or peptide 2[IL-8] are examples of agents falling under the scope of this invention which are useful for the prevention and treatment of malaria. Peptide 2, peptide 2 variants and derivatives are proinflammatory agents. Thus, these agents are useful to augment inflammatory responses, in particular weak inflammatory responses which are often associated with persistent infections such as parasitic infection, e.g., intracellular parasites.

5 Psoriasis. Psoriasis is an inflammatory disorder that is associated with MCP-1 and monocyte recruitment. Topical application of a therapeutic agent of the invention, e.g., peptide 3, is preferred to prevent or treat psoriasis as this delivery method reduces bioavailability problems. Derivatives of the therapeutic agents of the invention, e.g., CRD peptides, which are administered topically may exhibit enhanced bioavailability relative to non-derivatized counterparts.

10 Autoimmune Diseases. Autoimmune diseases, such as multiple sclerosis, Crohn's disease, rheumatoid arthritis and systemic lupus erythematosus, are characterized by inappropriate activation of the immune system, orchestrated by autoreactive leukocytes. Although it remains unclear what factors lead to the initial inappropriate recognition of self-antigens, a number of pro-inflammatory cytokines have been implicated in the continuing inflammation which underlies the tissue destruction that, in turn, leads to the morbidity and mortality associated with these 15 diseases. Of these inflammatory cytokines, TNF- α and the chemokines (in particular MIP-1 α) have been implicated.

20 For example, elevated MIP-1 α expression is detected in experimental autoimmune encephalomyelitis, a model of T-cell mediated autoimmune disease with some common characteristics to human multiple sclerosis. Elevated MIP1 α activity is also detected in the cerebrospinal fluid of patients with multiple sclerosis. Antibody therapy to reduce chemokine levels has been shown to be effective in animal models of autoimmune diseases, but this method only lowers chemokine 25 levels for a short period, and is unlikely to be useful in human therapy. In contrast, a general antagonist of chemokine signaling is likely to suppress the inappropriate

inflammation indefinitely. Thus, peptide 3, its derivatives and variants, may be useful to prevent and/or treat autoimmune disorders including, but not limited to, type I diabetes, multiple sclerosis, rheumatoid arthritis and systemic lupus erythematosus.

5 Moreover, different chemokine expression patterns may be associated with different autoimmune disorders, and hence each autoimmune disease may require a different derivative or variant of peptide 3. For example, MIP1 α may play a central role in multiple sclerosis. MIP1 α is a CC chemokine. Thus, the administration of a CC-selective agent of the invention can be used to treat multiple sclerosis (e.g.,

10 Leu₄Ile₁₁peptide 3(1-12)[MCP-1] or Ser₇Glu₈Glu₉peptide 3(1-12)[MCP-1]).

Wound Healing. Following wounding, there is a complex process of wound healing involving recruitment and proliferation of different cell types, elaboration of matrix, and increased immune surveillance. In the fetus (where increased immune surveillance is not required) this wound healing process leads to complete

15 restoration of the normal tissue architecture (e.g., normal dermal architecture is restored after incisional wounding). In marked contrast, in the adult, incisional wounding results in a wound healing process that does not restore normal dermal architecture. Matrix is elaborated in excess amounts and in inappropriate spatial organization. The result is a scar. In some cases, such as in children following

20 severe wounding such as from burns, the scars are hypertrophic having huge excess of matrix deposition and are particularly disfiguring.

 In adults, the risk of infection following wounding is high. Leukocytes, particularly neutrophils, are recruited rapidly to the wound site, while monocyte/macrophages appear several days after wounding, resulting in a rapid

25 formation of granulomatous tissue. Studies with antibodies have suggested that CXC chemokines such as IL-8 play an important role in neutrophil attraction to the wound site, and that inhibition of IL-8 production reduces both neutrophil accumulation and subsequent scarring. Experiments blocking CC chemokines have similarly shown that they have a role in the attraction of macrophages to the wound

site, and these cells may also promote rapid healing at the expense of wound quality. Hence inhibition of either CXC or CC chemokines, or both, may result in a decrease in the wound-induced inflammatory reaction, and in turn promote a balance between fast healing and good restoration of dermal architecture.

5 To prevent or reduce scarring and/or enhance wound healing, a preferred embodiment of the invention is the topical application of a therapeutic agent of the invention that inhibits chemokine action at the site of the wound. Thus, a broad spectrum chemokine inhibitor, such as peptide 3(1-12)[MCP-1], Leu₄Ile₁₁peptide 3(1-12)[MCP-1], CRD-Leu₄-Ile₁₁peptide 3[MCP-1] or WVQ, or combinations thereof may be administered. Alternatively, a selective inhibitor of IL-8, such as KEN, or a selective inhibitor of MCP-1, such as KQK, as well as combinations thereof may be administered. In addition, a combination of a broad spectrum inhibitor and a selective inhibitor may be administered. In this way, the various components of the wound-induced inflammatory process may be controlled as 10 desired and the wound may be allowed to heal more slowly (under conditions where it is protected from infection, e.g., by simultaneous use of antibiotics) but with enhanced recovery of dermal architecture. See U.S. Patent No. 5,202,118 for 15 methods to determine the efficacy of an agent to treat or enhance wound healing.

Hypertension. Hypertension is a risk factor for atherosclerosis. To determine 20 whether an agent of the invention is useful to inhibit or treat hypertension, a rabbit model is employed. New Zealand white rabbits are fed an atherogenic diet for three weeks to induce plaque formation. One half of each group of rabbits is administered an agent of the invention. Aortic coarctation is created in one group of the rabbits by wrapping a Dacron band around the midportion of the descending thoracic aorta 25 (stenosis group). Another group of rabbits undergo the banding technique without aortic constriction. Yet another group of rabbits serve as nonoperated controls. Monocyte binding to the aortic endothelial surface is determined with epifluorescent microscopy on standard aortic segments proximal and distal to the band. Immunohistochemistry is performed using the following antibodies:VCAM-1,

RAM11, CD11b, and factor VIII. In rabbits that did not receive the agent, hypertensive regions of the aorta proximal to the stenosis, monocyte adhesion and endothelial VCAM-1 expression are increased, with intimal thickening and accumulation of macrophage. In agent-treated rabbits, monocyte adhesion and 5 endothelial VCAM-1 expression, intimal thickening and accumulation of macrophage are decreased relative to non-agent-treated rabbits.

Tuberculosis. Infection with *Mycobacterium tuberculosis* is an example of a disease where the monocyte/macrophage is responsible for attempting to clear the pathogen from the host tissue (in this case the lung) but where the immune response 10 is often insufficient. As a consequence, even when antibiotics are used, *M. tuberculosis* infection can persist, clinically or sub-clinically as the body fails to clear the entire pathogen load.

Thus, diseases such as tuberculosis are amenable to therapy by agents which augment the existing immune reaction to recruit additional monocytes to the target 15 tissue. Agents which illicit a systemic inflammation are less useful because the systemic inflammation has side effects (e.g., nausea, high temperature, lethargy, etc.). Thus, agents which bind to DARC with high affinity, may be useful as these agents may augment an existing inflammatory reaction but do not result in systemic inflammation. Since DARC normally acts to limit the extent of a local 20 inflammatory reaction by sequestering locally produced chemokines, agents which reduce or block the capacity for DARC to sequester the chemokines may augment the desired response. Thus, one embodiment of the invention is the administration of a DARC binding agent, e.g., peptide 2[MCP-1], either locally to the lungs or systemically, to reduce or inhibit DARC sequestration of chemokines. Increased 25 local chemokine levels can augment monocyte recruitment, and the resultant increase in the number of tissue macrophages may aid clearance of *M. tuberculosis* and reduce or prevent chronic infection.

The agent of the invention may be administered either alone, or more preferably, in combination with antibiotics or other drugs which have been shown to

inhibit the growth of *M. tuberculosis*, but which when used singly may not prevent or abolish chronic infection.

Agents such as peptide 2[MCP-1] which augment existing immune reactions may also be useful for reduction or elimination of other chronic or parasitic

5 infections, e.g., leishmania, trypanosomes, leprosy, and the like.

Basophil-mediated diseases. Asthma is a disease characterized by hyper-reactive airways and chronic inflammation resulting from an influx of many cell types and inflammatory mediators. The interaction and causal effects of all the inflammatory mediators in asthma is not entirely understood. MCP-1 can play a role

10 in asthma through several different effector functions such as: monocyte recruitment, basophil recruitment, lymphocyte recruitment, monocyte activation or by triggering the release of histamine from basophils or resident mast cells (Bischoff et al., *J. Exp. Med.*, 175(5), 1271 (1992)). Inhibition of these processes are likely to reduce the severity of the disease. Allergic diseases, like asthma, are manifested

15 through a complex interaction of inflammatory mediators including monocytes/macrophages, lymphocytes and histamine release from mast cells and basophils.

A preferred mode for administration of a therapeutic agent of the invention to treat or inhibit the symptoms associated with asthma is by inhalation. As red

20 blood cells are not normally present in the respiratory tract, the DARC specificity of the therapeutic agent is less important for administration to the respiratory tract than for other modes of administration.

Endotoxemia. Endotoxemia is an acute systemic illness often mediated by LPS, a major component in the cell wall of gram-negative bacteria. LPS stimulates

25 the release of proinflammatory cytokines. MCP-1 and MCP-2 are expressed in endotoxemia and exert their effect by recruiting leukocytes to target organs. The intraperitoneal administration of recombinant murine MCP-1 to LPS-challenged mice protected them from endotoxic lethality (Zisman et al., *J. Clin. Invest.*, 99,

2832 (1997)). Thus, preferred peptides for use in this embodiment of the invention are MCP-1 and MCP-2 peptides.

Myocardial Infarction/Acute Ischemia. Myocardial infarction is the result of acute closure of a coronary vessel usually due to thrombosis secondary to rupture of an atherosclerotic plaque. The damage to the adjacent myocardium and resultant heart failure is secondary to the period of ischemia and the damage caused during the reperfusion period. Reperfusion injuries are associated with increased oxygen free radicals and inflammatory mediators. MCP-1 is up-regulated during the reperfusion period and is a key inflammatory mediator (Kumar et al., *Circulation*, 10, 90, 1427 (1994); Kumar et al., *Circulation*, 95, 693 (1997)). Inhibition of MCP-1 and the resultant inflammatory input may decrease damage to the myocardium during recovery and reduce the incidence of heart failure.

Rheumatoid Arthritis. Rheumatoid arthritis is a multi-systemic inflammatory disease involving primarily the joints but also the skin, blood vessels, heart, lung and muscle. The characteristic pathology of rheumatoid arthritis involves the accumulation of non-suppurative inflammatory cell infiltrate consisting of macrophages and lymphocytes within the joint. MCP-1 is produced by both synovial cells and infiltrating monocyte/macrophages in rheumatoid arthritis and is thought to contribute to the accumulation of inflammatory cells within the joint. The ability of native MCP-1 and an antagonist of MCP-1 (residues 9-76 of native MCP-1) have been assessed in the MRL-lpr model of chronic arthritis. Treatment with the antagonist MCP-1 (9-76) but not native MCP-1 resulted in a reduction of the symptoms and histopathology of chronic arthritis in this model (Gong et al., *J. Exp. Med.*, 186, 131 (1997); Plater-Zyberk et al., *Immunol. Lett.*, 57, 117 (1997); Wilder, *Clin. Rheumat.*, 10, 259 (1996)). Thus, peptide 3, its variants, analogs and derivatives may be especially useful to treat or prevent rheumatoid arthritis.

Contraception. Knockout mice for the CXCR4 chemokine receptor exhibit embryonic lethality. Agents of the invention have been identified which block the CXCR4 receptor (see Example 5) and other chemokine receptors. Thus, the agents

of the invention may be useful in inducing abortion or providing contraception. Blockade of the CXCR4 receptor could provide an alternative to traditional contraceptives and could be used post-coitus.

IV. Dosages, Formulations and Routes of Administration of the Agents of the

5 Invention

The therapeutic agents of the invention, including a compound of formula (I)-(V), including their salts, are preferably administered so as to achieve serum levels of about 0.01 pM to about 100 nM, more preferably at doses of about 0.01 pM to about 5 nM, and even more preferably at doses of about 0.1 pM to about 10 2 nM, of the therapeutic agent. To achieve these levels, the agent may be administered at dosages of at least about 0.01 to about 100 mg/kg, more preferably about 0.1 to about 50 mg/kg, and even more preferably about 0.1 to about 30 mg/kg, of body weight, although other dosages may provide beneficial results. The amount administered will vary depending on various factors including, but not limited to, 15 the agent chosen, the disease, whether prevention or treatment is to be achieved, and if the agent is modified for bioavailability and *in vivo* stability.

Administration of sense or antisense nucleic acid molecule may be accomplished through the introduction of cells transformed with an expression cassette comprising the nucleic acid molecule (see, for example, WO 93/02556) or 20 the administration of the nucleic acid molecule (see, for example, Felgner et al., U.S. Patent No. 5,580,859, Pardoll et al., Immunity, 3, 165 (1995); Stevenson et al., Immunol. Rev., 145, 211 (1995); Molling, J. Mol. Med., 75, 242 (1997); Donnelly et al., Ann. N.Y. Acad. Sci., 772, 40 (1995); Yang et al., Mol. Med. Today, 2, 476 (1996); Abdallah et al., Biol. Cell, 85, 1 (1995)). Pharmaceutical formulations, 25 dosages and routes of administration for nucleic acids are generally disclosed, for example, in Felgner et al., *supra*.

The amount of therapeutic agent administered is selected to treat a particular indication. For example, to treat malaria, higher doses of peptide 2, its variants or derivatives, may be administered, while smaller doses of peptide 2, its variants or

derivatives, are useful to prevent or inhibit HIV infection. The therapeutic agents of the invention are also amenable to chronic use for prophylactic purposes, preferably by systemic administration.

Administration of the therapeutic agents in accordance with the present invention may be continuous or intermittent, depending, for example, upon the recipient's physiological condition, whether the purpose of the administration is therapeutic or prophylactic, and other factors known to skilled practitioners. The administration of the agents of the invention may be essentially continuous over a preselected period of time or may be in a series of spaced doses. Both local and systemic administration is contemplated.

One or more suitable unit dosage forms comprising the therapeutic agents of the invention, which, as discussed below, may optionally be formulated for sustained release, can be administered by a variety of routes including oral, or parenteral, including by rectal, buccal, vaginal and sublingual, transdermal, subcutaneous, intravenous, intramuscular, intraperitoneal, intrathoracic, intrapulmonary and intranasal routes. The formulations may, where appropriate, be conveniently presented in discrete unit dosage forms and may be prepared by any of the methods well known to pharmacy. Such methods may include the step of bringing into association the therapeutic agent with liquid carriers, solid matrices, semi-solid carriers, finely divided solid carriers or combinations thereof, and then, if necessary, introducing or shaping the product into the desired delivery system.

When the therapeutic agents of the invention are prepared for oral administration, they are preferably combined with a pharmaceutically acceptable carrier, diluent or excipient to form a pharmaceutical formulation, or unit dosage form. The total active ingredients in such formulations comprise from 0.1 to 99.9% by weight of the formulation. By "pharmaceutically acceptable" it is meant the carrier, diluent, excipient, and/or salt must be compatible with the other ingredients of the formulation, and not deleterious to the recipient thereof. The active ingredient for oral administration may be present as a powder or as granules; as a solution, a

suspension or an emulsion; or in achievable base such as a synthetic resin for ingestion of the active ingredients from a chewing gum. The active ingredient may also be presented as a bolus, electuary or paste.

Formulations suitable for vaginal administration may be presented as 5 pessaries, tampons, creams, gels, pastes, douches, lubricants, foams or sprays containing, in addition to the active ingredient, such carriers as are known in the art to be appropriate. Formulations suitable for rectal administration may be presented as suppositories.

Pharmaceutical formulations containing the therapeutic agents of the 10 invention can be prepared by procedures known in the art using well known and readily available ingredients. For example, the agent can be formulated with common excipients, diluents, or carriers, and formed into tablets, capsules, suspensions, powders, and the like. Examples of excipients, diluents, and carriers that are suitable for such formulations include the following fillers and extenders 15 such as starch, sugars, mannitol, and silicic derivatives; binding agents such as carboxymethyl cellulose, HPMC and other cellulose derivatives, alginates, gelatin, and polyvinyl-pyrrolidone; moisturizing agents such as glycerol; disintegrating agents such as calcium carbonate and sodium bicarbonate; agents for retarding dissolution such as paraffin; resorption accelerators such as quaternary ammonium 20 compounds; surface active agents such as cetyl alcohol, glycerol monostearate; adsorptive carriers such as kaolin and bentonite; and lubricants such as talc, calcium and magnesium stearate, and solid polyethyl glycols.

For example, tablets or caplets containing the agents of the invention can include buffering agents such as calcium carbonate, magnesium oxide and 25 magnesium carbonate. Caplets and tablets can also include inactive ingredients such as cellulose, pregelatinized starch, silicon dioxide, hydroxy propyl methyl cellulose, magnesium stearate, microcrystalline cellulose, starch, talc, titanium dioxide, benzoic acid, citric acid, corn starch, mineral oil, polypropylene glycol, sodium phosphate, and zinc stearate, and the like. Hard or soft gelatin capsules containing

an agent of the invention can contain inactive ingredients such as gelatin, microcrystalline cellulose, sodium lauryl sulfate, starch, talc, and titanium dioxide, and the like, as well as liquid vehicles such as polyethylene glycols (PEGs) and vegetable oil. Moreover, enteric coated caplets or tablets of an agent of the invention
5 are designed to resist disintegration in the stomach and dissolve in the more neutral to alkaline environment of the duodenum.

The therapeutic agents of the invention can also be formulated as elixirs or solutions for convenient oral administration or as solutions appropriate for parenteral administration, for instance by intramuscular, subcutaneous or
10 intravenous routes.

The pharmaceutical formulations of the therapeutic agents of the invention can also take the form of an aqueous or anhydrous solution or dispersion, or alternatively the form of an emulsion or suspension.

Thus, the therapeutic agent may be formulated for parenteral administration
15 (e.g., by injection, for example, bolus injection or continuous infusion) and may be presented in unit dose form in ampules, pre-filled syringes, small volume infusion containers or in multi-dose containers with an added preservative. The active ingredients may take such forms as suspensions, solutions, or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending,
20 stabilizing and/or dispersing agents. Alternatively, the active ingredients may be in powder form, obtained by aseptic isolation of sterile solid or by lyophilization from solution, for constitution with a suitable vehicle, e.g., sterile, pyrogen-free water, before use.

These formulations can contain pharmaceutically acceptable vehicles and
25 adjuvants which are well known in the prior art. It is possible, for example, to prepare solutions using one or more organic solvent(s) that is/are acceptable from the physiological standpoint, chosen, in addition to water, from solvents such as acetone, ethanol, isopropyl alcohol, glycol ethers such as the products sold under the name "Dowanol", polyglycols and polyethylene glycols, C₁-C₄ alkyl esters of short-

chain acids, preferably ethyl or isopropyl lactate, fatty acid triglycerides such as the products marketed under the name "Miglyol", isopropyl myristate, animal, mineral and vegetable oils and polysiloxanes.

The compositions according to the invention can also contain thickening agents such as cellulose and/or cellulose derivatives. They can also contain gums such as xanthan, guar or carbo gum or gum arabic, or alternatively polyethylene glycols, bentones and montmorillonites, and the like.

It is possible to add, if necessary, an adjuvant chosen from antioxidants, surfactants, other preservatives, film-forming, keratolytic or comedolytic agents, perfumes and colorings. Also, other active ingredients may be added, whether for the conditions described or some other condition.

For example, among antioxidants, t-butylhydroquinone, butylated hydroxyanisole, butylated hydroxytoluene and α -tocopherol and its derivatives may be mentioned. The galenical forms chiefly conditioned for topical application take the form of creams, milks, gels, dispersion or microemulsions, lotions thickened to a greater or lesser extent, impregnated pads, ointments or sticks, or alternatively the form of aerosol formulations in spray or foam form or alternatively in the form of a cake of soap.

Additionally, the agents are well suited to formulation as sustained release dosage forms and the like. The formulations can be so constituted that they release the active ingredient only or preferably in a particular part of the intestinal or respiratory tract, possibly over a period of time. The coatings, envelopes, and protective matrices may be made, for example, from polymeric substances, such as polylactide-glycolates, liposomes, microemulsions, microparticles, nanoparticles, or waxes. These coatings, envelopes, and protective matrices are useful to coat indwelling devices, e.g., stents, catheters, peritoneal dialysis tubing, and the like.

The therapeutic agents of the invention can be delivered via patches for transdermal administration. See U.S. Patent No. 5,560,922 for examples of patches suitable for transdermal delivery of a therapeutic agent. Patches for transdermal

delivery can comprise a backing layer and a polymer matrix which has dispersed or dissolved therein a therapeutic agent, along with one or more skin permeation enhancers. The backing layer can be made of any suitable material which is impermeable to the therapeutic agent. The backing layer serves as a protective

5 cover for the matrix layer and provides also a support function. The backing can be formed so that it is essentially the same size layer as the polymer matrix or it can be of larger dimension so that it can extend beyond the side of the polymer matrix or overlay the side or sides of the polymer matrix and then can extend outwardly in a manner that the surface of the extension of the backing layer can be the base for an

10 adhesive means. Alternatively, the polymer matrix can contain, or be formulated of, an adhesive polymer, such as polyacrylate or acrylate/vinyl acetate copolymer. For long-term applications it might be desirable to use microporous and/or breathable backing laminates, so hydration or maceration of the skin can be minimized.

Examples of materials suitable for making the backing layer are films of high

15 and low density polyethylene, polypropylene, polyurethane, polyvinylchloride, polyesters such as poly(ethylene phthalate), metal foils, metal foil laminates of such suitable polymer films, and the like. Preferably, the materials used for the backing layer are laminates of such polymer films with a metal foil such as aluminum foil. In such laminates, a polymer film of the laminate will usually be in contact with the

20 adhesive polymer matrix.

The backing layer can be any appropriate thickness which will provide the desired protective and support functions. A suitable thickness will be from about 10 to about 200 microns.

Generally, those polymers used to form the biologically acceptable adhesive

25 polymer layer are those capable of forming shaped bodies, thin walls or coatings through which therapeutic agents can pass at a controlled rate. Suitable polymers are biologically and pharmaceutically compatible, nonallergenic and insoluble in and compatible with body fluids or tissues with which the device is contacted. The use of soluble polymers is to be avoided since dissolution or erosion of the matrix by

skin moisture would affect the release rate of the therapeutic agents as well as the capability of the dosage unit to remain in place for convenience of removal.

Exemplary materials for fabricating the adhesive polymer layer include polyethylene, polypropylene, polyurethane, ethylene/propylene copolymers, 5 ethylene/ethylacrylate copolymers, ethylene/vinyl acetate copolymers, silicone elastomers, especially the medical-grade polydimethylsiloxanes, neoprene rubber, polyisobutylene, polyacrylates, chlorinated polyethylene, polyvinyl chloride, vinyl chloride-vinyl acetate copolymer, crosslinked polymethacrylate polymers (hydro-gel), polyvinylidene chloride, poly(ethylene terephthalate), butyl rubber, 10 epichlorohydrin rubbers, ethylenvinyl alcohol copolymers, ethylene-vinyloxyethanol copolymers; silicone copolymers, for example, polysiloxane-polycarbonate copolymers, polysiloxanepolyethylene oxide copolymers, polysiloxane-polymethacrylate copolymers, polysiloxane-alkylene copolymers (e.g., polysiloxane-ethylene copolymers), polysiloxane-alkylenesilane copolymers (e.g., polysiloxane-15 ethylenesilane copolymers), and the like; cellulose polymers, for example methyl or ethyl cellulose, hydroxy propyl methyl cellulose, and cellulose esters; polycarbonates; polytetrafluoroethylene; and the like.

Preferably, a biologically acceptable adhesive polymer matrix should be selected from polymers with glass transition temperatures below room temperature.

20 The polymer may, but need not necessarily, have a degree of crystallinity at room temperature. Cross-linking monomeric units or sites can be incorporated into such polymers. For example, cross-linking monomers can be incorporated into polyacrylate polymers, which provide sites for cross-linking the matrix after dispersing the therapeutic agent into the polymer. Known cross-linking monomers 25 for polyacrylate polymers include polymethacrylic esters of polyols such as butylene diacrylate and dimethacrylate, trimethylol propane trimethacrylate and the like. Other monomers which provide such sites include allyl acrylate, allyl methacrylate, diallyl maleate and the like.

Preferably, a plasticizer and/or humectant is dispersed within the adhesive polymer matrix. Water-soluble polyols are generally suitable for this purpose. Incorporation of a humectant in the formulation allows the dosage unit to absorb moisture on the surface of skin which in turn helps to reduce skin irritation and to

5 prevent the adhesive polymer layer of the delivery system from failing.

Therapeutic agents released from a transdermal delivery system must be capable of penetrating each layer of skin. In order to increase the rate of permeation of a therapeutic agent, a transdermal drug delivery system must be able in particular to increase the permeability of the outermost layer of skin, the stratum corneum,

10 which provides the most resistance to the penetration of molecules. The fabrication of patches for transdermal delivery of therapeutic agents is well known to the art.

For administration to the upper (nasal) or lower respiratory tract by inhalation, the therapeutic agents of the invention are conveniently delivered from an insufflator, nebulizer or a pressurized pack or other convenient means of

15 delivering an aerosol spray. Pressurized packs may comprise a suitable propellant such as dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol, the dosage unit may be determined by providing a valve to deliver a metered amount.

Alternatively, for administration by inhalation or insufflation, the

20 composition may take the form of a dry powder, for example, a powder mix of the therapeutic agent and a suitable powder base such as lactose or starch. The powder composition may be presented in unit dosage form in, for example, capsules or cartridges, or, e.g., gelatine or blister packs from which the powder may be administered with the aid of an inhalator, insufflator or a metered-dose inhaler.

25 For intra-nasal administration, the therapeutic agent may be administered via nose drops, a liquid spray, such as via a plastic bottle atomizer or metered-dose inhaler. Typical of atomizers are the Mistometer (Wintrop) and the Medihaler (Riker).

The local delivery of the therapeutic agents of the invention can also be by a variety of techniques which administer the agent at or near the site of disease. Examples of site-specific or targeted local delivery techniques are not intended to be limiting but to be illustrative of the techniques available. Examples include local 5 delivery catheters, such as an infusion or indwelling catheter, e.g., a needle infusion catheter, shunts and stents or other implantable devices, site specific carriers, direct injection, or direct applications.

For topical administration, the therapeutic agents may be formulated as is known in the art for direct application to a target area. Conventional forms for this 10 purpose include wound dressings, coated bandages or other polymer coverings, ointments, creams, lotions, pastes, jellies, sprays, and aerosols, as well as in toothpaste and mouthwash, or by other suitable forms, e.g., via a coated condom. Ointments and creams may, for example, be formulated with an aqueous or oily base with the addition of suitable thickening and/or gelling agents. Lotions may be 15 formulated with an aqueous or oily base and will in general also contain one or more emulsifying agents, stabilizing agents, dispersing agents, suspending agents, thickening agents, or coloring agents. The active ingredients can also be delivered via iontophoresis, e.g., as disclosed in U.S. Patent Nos. 4,140,122; 4,383,529; or 4,051,842. The percent by weight of a therapeutic agent of the invention present in 20 a topical formulation will depend on various factors, but generally will be from 0.01% to 95% of the total weight of the formulation, and typically 0.1-25% by weight.

When desired, the above-described formulations can be adapted to give sustained release of the active ingredient employed, e.g., by combination with 25 certain hydrophilic polymer matrices, e.g., comprising natural gels, synthetic polymer gels or mixtures thereof.

Drops, such as eye drops or nose drops, may be formulated with an aqueous or non-aqueous base also comprising one or more dispersing agents, solubilizing agents or suspending agents. Liquid sprays are conveniently delivered from

pressurized packs. Drops can be delivered via a simple eye dropper-capped bottle, or via a plastic bottle adapted to deliver liquid contents dropwise, via a specially shaped closure.

The therapeutic agent may further be formulated for topical administration in 5 the mouth or throat. For example, the active ingredients may be formulated as a lozenge further comprising a flavored base, usually sucrose and acacia or tragacanth; pastilles comprising the composition in an inert base such as gelatin and glycerin or sucrose and acacia; mouthwashes comprising the composition of the present invention in a suitable liquid carrier; and pastes and gels, e.g., toothpastes or gels, 10 comprising the composition of the invention.

The formulations and compositions described herein may also contain other ingredients such as antimicrobial agents, or preservatives. Furthermore, the active ingredients may also be used in combination with other therapeutic agents, for example, oral contraceptives, bronchodilators, anti-viral agents, steroids and the 15 like.

Sustained Released Dosage Forms

Sustained release dosage forms of the invention may comprise microparticles and/or nanoparticles having a therapeutic agent dispersed therein. The therapeutic dosage forms of this aspect of the present invention may be of any configuration 20 suitable for sustained release. Preferred sustained release therapeutic dosage forms exhibit one or more of the following characteristics:

- microparticles (e.g., from about 0.5 micrometers to about 100 micrometers in diameter, with about 0.5 to about 2 micrometers more preferred; or from about 0.01 micrometers to about 200 micrometers in diameter, preferably from about 0.5 25 to about 50 micrometers, and more preferably from about 2 to about 15 micrometers) or nanoparticles (e.g., from about 1.0 nanometer to about 1000 nanometers in diameter, with about 50 to about 250 nanometers being more preferred; or from about 0.01 nanometer to about 1000 nanometers in diameter, preferably from about 50 to about 200 nanometers), free flowing powder structure;

- biodegradable structure designed to biodegrade over a period of time preferably between from about 0.5 to about 180 days, preferably from about 1-3 to about 150 days, or from about 3 to about 180 days, with from about 10 to about 21 days more preferred; or non-biodegradable structure to allow therapeutic agent diffusion to occur over a time period of between from about 0.5 to about 180 days, more preferably from about 30 to about 120 days; or from about 3 to about 180 days, with from about 10 to about 21 days preferred;
- biocompatible with target tissue and the local physiological environment into which the dosage form to be administered, including yielding biocompatible biodegradation products;
- facilitate a stable and reproducible dispersion of therapeutic agent therein, preferably to form a therapeutic agent-polymer matrix, with active therapeutic agent release occurring by one or both of the following routes: (1) diffusion of the therapeutic agent through the dosage form (when the therapeutic agent is soluble in the shaped polymer or polymer mixture defining the dimensions of the dosage form); or (2) release of the therapeutic agent as the dosage form biodegrades; and/or
- for targeted dosage forms, capability to have, preferably, from about 1 to about 10,000 binding protein/peptide to dosage form bonds and more preferably, a maximum of about 1 binding peptide to dosage form bond per 150 square angstroms of particle surface area. The total number of binding protein/peptide to dosage form bonds depends upon the particle size used. The binding proteins or peptides are capable of coupling to the particles of the therapeutic dosage form through covalent ligand sandwich or non-covalent modalities as set forth herein.

Nanoparticle sustained release therapeutic dosage forms are preferably biodegradable and, optionally, bind to the vascular smooth muscle cells and enter those cells, primarily by endocytosis. The biodegradation of the nanoparticles occurs over time (e.g., 30 to 120 days; or 10 to 21 days) in prelysosomal vesicles and lysosomes. Preferred larger microparticle therapeutic dosage forms of the present invention release the therapeutic agents for subsequent target cell uptake

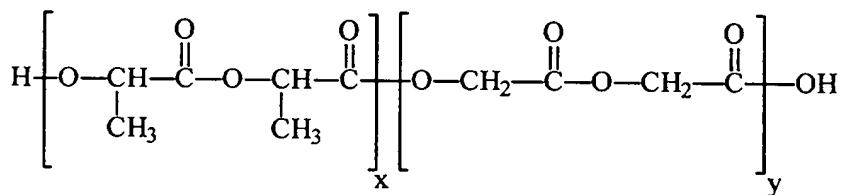
with only a few of the smaller microparticles entering the cell by phagocytosis. A practitioner in the art will appreciate that the precise mechanism by which a target cell assimilates and metabolizes a dosage form of the present invention depends on the morphology, physiology and metabolic processes of those cells. The size of the

5 particle sustained release therapeutic dosage forms is also important with respect to the mode of cellular assimilation. For example, the smaller nanoparticles can flow with the interstitial fluid between cells and penetrate the infused tissue. The larger microparticles tend to be more easily trapped interstitially in the infused primary tissue, and thus are useful to therapeutic agents.

10 Preferred sustained release dosage forms of the present invention comprise biodegradable microparticles or nanoparticles. More preferably, biodegradable microparticles or nanoparticles are formed of a polymer containing matrix that biodegrades by random, nonenzymatic, hydrolytic scissioning to release therapeutic agent, thereby forming pores within the particulate structure.

15 Polymers derived from the condensation of alpha hydroxycarboxylic acids and related lactones are preferred for use in the present invention. A particularly preferred moiety is formed of a mixture of thermoplastic polyesters (e.g., polylactide or polyglycolide) or a copolymer of lactide and glycolide components, such as poly(lactide-co-glycolide). An exemplary structure, a random poly(DL-lactide-co-

20 glycolide), is shown below, with the values of x and y being manipulable by a practitioner in the art to achieve desirable microparticle or nanoparticle properties.



Other agents suitable for forming particulate dosage forms of the present invention include polyorthoesters and polyacetals (Polymer Letters, 18:293 (1980) and polyorthocarbonates (U.S. Patent No. 4,093,709) and the like.

Preferred lactic acid/glycolic acid polymer containing matrix particles of the present invention are prepared by emulsion-based processes, that constitute modified solvent extraction processes, see, for example, processes described by Cowsar et al., "Poly(Lactide-Co-Glycolide) Microcapsules for Controlled Release of Steroids," Methods Enzymology, 112:101-116, 1985 (steroid entrapment in microparticles); Eldridge et al., "Biodegradable and Biocompatible Poly(DL-Lactide-Co-Glycolide) Microspheres as an Adjuvant for Staphylococcal Enterotoxin B Toxoid Which Enhances the Level of Toxin-Neutralizing Antibodies," Infection and Immunity, 59:2978-2986, 1991 (toxoid entrapment); Cohen et al., "Controlled Delivery Systems for Proteins Based on Poly(Lactic/Glycolic Acid) Microspheres," Pharmaceutical Research, 8(6):713-720, 1991 (enzyme entrapment); and Sanders et al., "Controlled Release of a Luteinizing Hormone-Releasing Hormone Analogue from Poly(D,L-Lactide-Co-Glycolide) Microspheres," J. Pharmaceutical Science, 73(9):1294-1297, 1984 (peptide entrapment).

In general, the procedure for forming particle dosage forms of the present invention involves dissolving the polymer in a halogenated hydrocarbon solvent, dispersing a therapeutic agent solution (preferably aqueous) therein, and adding an additional agent that acts as a solvent for the halogenated hydrocarbon solvent but not for the polymer. The polymer precipitates out from the polymer-halogenated hydrocarbon solution onto droplets of the therapeutic agent containing solution and entraps the therapeutic agent. Preferably the therapeutic agent is substantially uniformly dispersed within the sustained release dosage form of the present invention. Following particle formation, they are washed and hardened with an organic solvent. Water washing and aqueous nonionic surfactant washing steps follow, prior to drying at room temperature under vacuum.

For biocompatibility purposes, particulate dosage forms, characterized by a therapeutic agent dispersed in the matrix of the particles, are sterilized prior to packaging, storage or administration. Sterilization may be conducted in any convenient manner therefor. For example, the particles can be irradiated with 5 gamma radiation, provided that exposure to such radiation does not adversely impact the structure or function of the therapeutic agent dispersed in the therapeutic agent-polymer matrix or the binding protein/peptide attached thereto. If the therapeutic agent or binding protein/peptide is so adversely impacted, the particle dosage forms can be produced under sterile conditions.

10 Release of the therapeutic agent from the particle dosage forms of the present invention can occur as a result of both diffusion and particle matrix erosion. The biodegradation rate directly effects the kinetics of therapeutic agent release. The biodegradation rate is regulable by alteration of the composition or structure of the sustained release dosage form. For example, alteration of the lactide/glycolide ratio 15 in preferred dosage forms of the present invention can be conducted, as described by Tice et al., "Biodegradable Controlled-Release Parenteral Systems," Pharmaceutical Technology, pp. 26-35, 1984; by inclusion of agents that alter the rate of polymer hydrolysis, such as citric acid and sodium carbonate, as described by Kent et al., "Microencapsulation of Water Soluble Active Polypeptides," U.S. Patent No. 20 4,675,189; by altering the loading of therapeutic agent in the lactide/glycolide polymer, the degradation rate being inversely proportional to the amount of therapeutic agent contained therein, by judicious selection of an appropriate analog of a common family of therapeutic agents that exhibit different potencies so as to alter said core loadings; and by variation of particle size, as described by Beck et al., 25 "Poly(DL-Lactide-Co-Glycolide)/Norethisterone Microcapsules: An Injectable Biodegradable Contraceptive," Biol. Reprod., 28:186-195, 1983, or the like. All of the aforementioned methods of regulating biodegradation rate influence the intrinsic viscosity of the polymer containing matrix, thereby altering the hydration rate thereof.

The preferred lactide/glycolide structure is biocompatible with the mammalian physiological environment. Also, these preferred sustained release dosage forms have the advantage that biodegradation thereof forms lactic acid and glycolic acid, both normal metabolic products of mammals.

5 Functional groups required for binding of the protein/peptide to the particle dosage form are optionally included in or on the particle matrix and are attached to the non-degradable or biodegradable polymeric units. Functional groups that are useful for this purpose include those that are reactive with peptides, e.g., carboxyl groups, amine groups, sulfhydryl groups and the like. Preferred binding

10 enhancement moieties include the terminal carboxyl groups of the preferred (lactide-glycolide) polymer containing matrix or the like.

To employ the therapeutic agents of the invention to enhance the immunological response of a particular immunogen, e.g., the *Haemophilis influenza* type b (Hib) capsular polysaccharide (polyribosylribitol phosphate, PRP), the agents 15 may be conjugated to the immunogen. Thus, for example, peptide 2(1-15)[MCP-1] may be covalently linked to PRP through a 6 carbon spacer molecule derived from adipic acid dihydrazide (see Gordon, Patent 83/4939, Republic of South Africa, 1984), and administered in a manner similar to that described in Eskola et al., *Lancet*, 1, 1184 (1985). The use of nucleic acid molecules to prepare vaccines is 20 described in, for example, Felgner et al., *supra* and Stevenson et al., *supra*.

A vaccine of the invention may also comprise cells or viruses having nucleic acid encoding the immunogen and a peptide or variant peptide of the invention or its complement, optionally as a fusion protein.

For a general description of vaccine principles and practice, see Ada, In: 25 Fundamental Immunology, 2nd ed., Raven Press Ltd., N.Y., pp. 985-1030 (1989).

V. Detection of the Peptides of the Invention in Physiological Fluid

Analysis of peptide 3 in blood and urine was performed on a semi-permeable surface (SPS) HPLC column (restricted access media). Serum or other protein-containing samples can be injected directly onto an SPS column (e.g., SPS-C18 with

a column size of 4.6 mm × 250 mm; using a mobile phase: A: 0.1% TFA in water, B: 0.1% TFA in acetonitrile: 0-5 min – 5% B, 5-30 min – 60% B, 30-40 min – 5% B detector; 215 nm). The outer phase of the column forms a semipermeable surface that prevents large molecules from reaching the inner phase. Small molecules 5 penetrate the semipermeable surface and interact with the inner reversed phase.

Standards of peptide 3 (range of 1.5 µg/ml to 1000 µg/ml) in PBS were injected and a standard curve was created. 20 µl of serum and urine were injected and the areas under the peptide 3 peaks were obtained. The concentration was then calculated from the standard curve. This method can detect at least about 20 µg/ml 10 of a peptide in physiological fluid samples.

The invention will be further described by, but is not limited to, the following examples.

Example 1

15 Identification and Characterization of Pan-Chemokine Peptide Inhibitors

Based on an alignment of MCP-1 sequences from different species, three regions in MCP-1 were identified which were conserved between all the species examined. Three purified (>95% purity) peptides (12-15mers) were prepared which had the greatest sequence homology between the human and mouse MCP-1 20 sequences (Table 1). These peptides were screened for their ability to inhibit hMCP-1 induced THP-1 migration. Similarly, the sequences of *Xenopus laevis* TGF-beta1 and TGF-beta3 and human TGF-beta1 and TGF-beta3 were compared, and 3 regions (each 10mer) of perfect homology were identified. These peptides were termed “betatides”.

25 For this assay, THP-1 cells were maintained at a density of 4×10^5 cells per ml in RPMI-1640 supplemented with 10% fetal calf serum + 20 µM 2-mercaptoethanol. Chemotaxis was induced in a 96-well disposable chemotaxis chamber fitted with a 5 µM polycarbonate filter (PVP free, ChemoTX, Neuroprobe Inc., Cabin John). Twenty-nine µl of chemoattractant (recombinant human

chemokine; 50 ng/ml, i.e., 5.9 nM) or control (100 ng/ml TGF β) was added to the lower compartment of each well. The framed filter was aligned with the holes in the corner of the filter frame and placed over the wells. Five \times 10⁴ THP-1 cells in 25 μ l of RPMI-1640 culture media were added to the upper compartment. Peptides were 5 dissolved in Milli Q water and then serially diluted in culture medium. In most cases, the serially diluted peptides were added to the upper compartment of the chemotaxis chamber. The chamber was incubated at 37°C in a humidified atmosphere of 5% CO₂ for 4 hours.

After incubation, the cells were gently removed from the top of the filter 10 with a pipette, 20 μ l of 20 mM EDTA in PBS was added into each top well, and the mixture was incubated for 20 minutes at 4°C. The filter was then carefully flushed with media using a gentle flow, and removed. A standard curve was prepared to accurately quantify the number of THP-1 cells that had migrated. The curve was based on a two-fold dilution series of THP-1 cells (top standard 100,000 cells in 15 29 μ l). Cells which had migrated, and in separate wells, the cells in the standards, were stained with 3 μ l of MTT stock solution which was added directly into each well (5 mg/ml in RPMI 1640 without phenol red, Sigma Chemical Co.) and incubated at 37°C for 4 hours. The media was carefully aspirated from each well, and the converted dye was solubilized by 20 μ l of DMSO. Absorbance of converted 20 dye was measured at a wavelength of 595 nM using an ELISA plate reader. The number of cells that had migrated in each well was determined by interpolation of the standard curve.

Peptide 1[MCP-1] (see Table 1; SEQ ID NO:2), i.e., the N-terminal peptide 25 of human MCP-1, was only weakly active in the migration assay (ED₅₀ > 100 μ M; 10% inhibition at 100 μ M, p = 0.27). Peptide 2[MCP-1] (Table 1; SEQ ID NO:3) was also a weak inhibitor of chemokine-induced migration (ED₅₀ > 100 μ M; 19% inhibition at 100 μ M, p = 0.09). Thus, in the presence of a strong agonist, i.e., MCP-1, peptide 2[MCP-1] having SEQ ID NO:3, a weak agonist, displaces MCP-1 from its receptor. However, in the absence of a strong agonist, i.e., MCP-1, peptide

2[MCP-1] exhibited weak agonist properties, i.e., peptide 2[MCP-1] stimulated chemotaxis. Surprisingly, peptide 2(1-15)[SDF1 α] had potent pan-chemokine antagonist properties.

In contrast, peptide 3(1-12)[MCP-1] (Table 1; SEQ ID NO:1) was a highly effective inhibitor of MCP-1 induced THP-1 migration with a dose giving 50% inhibition (ED_{50}) of $8 \pm 1 \mu\text{M}$ (n=4). A typical dose response curve is shown in Figure 2. At concentrations above 50 μM , peptide 3(1-12)[MCP-1] having SEQ ID NO:1 abolished all of the MCP-1 induced THP-1 migration.

TABLE 1 Alignment of Chemokine Sequences

		Peptide 1		Peptide 2		Peptide 3	
AQPDAINAFLV	TCCYNTVNKRK	IPIQRLESYR	RITNIQCPKE	AVIFKTKRGK	EVCADPKERW	VRSNKHLDQ	IFQNLKP
AQPVGINTST	TCCYRFINKK	IPKQRLESYR	RTTSSHCPRE	AVIFKTKLDK	EICADPTQKW	VQDFMKHLDK	KTQTPKL
SASLAAADPTT	ACCFSYTTSRQ	IPQNFIADYF	E-TSSQCSKP	GVIFLTKRSR	QVCADPSEEW	VQKYVSDLEL	SA
SAPMGSDPPT	ACCFSYTARK	I,PRNFVVDYY	E-TSSLCSQP	AVVFQTKRSK	QVCADPSEW	VQEYYVYDLEL	N
HPGIPS	ACCFSYTARK	I,PRNFVVDYY	E-TSSLCSQP	AVVFQTKRSK	QVCADPSEW	VQEYYVYDLEL	N
LPRSAKELRC	QCIKTYSKPF	HPKFKEILRV	IESGPHCANT	EIIVRLSDGR	ELCLDPKENW	VQRVEKFLKR	AENS
GKPVSLSY	RCPCKRFRESH	IARANVKHLK	ILNTPNCALQ	IVARLKNNNR	QVCIDPKLKW	IQEYEKALNK	
		C...F...	I.....I	T...C...	AVI.....K	WVQ.....L...	...CONSENSUS

To determine whether the peptides were MCP-1 receptor antagonists, the peptides were introduced with the chemokine in the lower compartment (as opposed to with the cells in the upper compartment in the experiments described above; in the trans-well THP-1 migration assay. Under these conditions, peptide 1[MCP-1] having SEQ ID NO:2 was a more efficient inhibitor of MCP-1 induced chemokine migration than it had been when it was incubated with the cells, inhibiting 48% of the MCP-1 induced migration at 100 μ M compared to 10% inhibition when peptide 1[MCP-1] (SEQ ID NO:2) was incubated with the cells. This result is consistent with published reports which show that peptide 1[MCP-1] (SEQ ID NO:2) and its derivatives act by disrupting the MCP-1 dimer, forming inactive monomers. Peptide 1[MCP-1] (SEQ ID NO:2) is not, therefore, a classical receptor-level antagonist of MCP-1 function. In marked contrast, peptide 3(1-12)[MCP-1] having SEQ ID NO:1 was much less effective when incubated with the chemokine than with the cells (17% inhibition at 100 μ M compared with >99% inhibition), suggesting that a peptide having SEQ ID NO:1 inhibits MCP-1 induced migration by direct receptor antagonism. To confirm this observation, the binding affinity of an N-terminally biotinylated derivative of peptide 3(1-12)[MCP-1] (SEQ ID NO:1) was determined. This derivative bound to the surface of THP-1 cells with a k_a of about 10 μ M.

Peptide 3(1-12)[MCP-1] (SEQ ID NO:1) also inhibited other functions of MCP-1, which may be mediated by different combinations of receptors. MCP-1 has been reported to be a weak co-mitogen with 0.5% fetal calf serum for cultured smooth muscle cells. It was found that 100 μ M peptide 3(1-12)[MCP-1] (SEQ ID NO:1) completely abolished the co-mitogenic effect of MCP-1 for cultured smooth muscle cells, also consistent with the hypothesis that peptide 3(1-12)[MCP-1] (SEQ ID NO:1) is an MCP-1 receptor antagonist. The observation that peptide 3(1-12)[MCP-1] (SEQ ID NO:1) completely inhibits different responses to MCP-1 in different cell types suggests that peptide 3 may be a general antagonist of all chemokine receptors capable of binding and signaling in response to MCP-1.

To investigate the receptor specificity of peptide 3 inhibition, the ED₅₀ was determined for peptide 3(1-12)[MCP-1] (SEQ ID NO:1) inhibition of THP-1 migration induced by chemokines which signal through different receptors than MCP-1 receptors. Representative chemokines included a beta-chemokine ("CC"), 5 MIP-1 α , and two alpha-chemokines ("CXC"), IL-8 and SDF-1 α . Additionally, to determine the specificity of peptide 3(1-12)[MCP-1] (SEQ ID NO:1) for chemokine receptors, TGF-beta was selected as a migration-inducing agent unrelated to the chemokine family, and as an agent which elicits a biological activity by signaling through identified, unrelated receptors.

10 Peptide 3(1-12)[MCP-1] (SEQ ID NO:1) inhibited the THP-1 migration induced response to all four of the selected chemokines, with the order of potency: MIP-1 α \geq MCP-1 $>$ SDF1 α \geq IL-8 (see Table 2). In contrast, peptide 1[MCP-1] (SEQ ID NO:2) or peptide 2(1-15)[MCP-1] (SEQ ID NO:3) did not inhibit migration in response to any of these chemokines by more than 20%, even at 15 100 μ M (Table 2).

TABLE 2**(a) ED₅₀ for inhibition of THP-1 migration**

5	PEPTIDE	ED ₅₀ (μM) versus				
		MCP-1	MIP1 α	IL8	SDF-1 α	TGF β 1
	Peptide 1 (SEQ ID NO:2)	n.s. ^b	n.s. ^b	n.s.	n.s.	n.s.
	Peptide 2 (SEQ ID NO:3)	n.s.	n.s.	n.s.	n.s.	n.s.
10	Peptide 3 ^a (SEQ ID NO:1)	8 ± 1	8 ± 1	14 ± 1	10 ± 0	n.s.

(b) Extent of inhibition of THP-1 migration at 100 μM

15	PEPTIDE	% inhibition at 100 μM versus				
		MCP-1	MIP1 α	IL8	SDF-1 α	TGF β 1
	Peptide 1 (SEQ ID NO:2)	n.s. ^b	n.s. ^b	n.s.	n.s.	n.s.
	Peptide 2 (SEQ ID NO:3)	n.s.	n.s.	n.s.	n.s.	n.s.
20	Peptide 3 (SEQ ID NO:1)	112	99	103	107	n.s.

(c) Extent of inhibition of human monocyte migration at 100 μM

25	PEPTIDE	% inhibition at 100 μM versus				
		MCP-1	MIP1 α	IL8	SDF-1 α	TGF β 1
	Peptide 1 (SEQ ID NO:2)	n.s.	n.s.	n.s.	n.s.	n.s.
	Peptide 2 (SEQ ID NO:3)	23	n.s.	n.s.	n.s.	n.s.
30	Peptide 3 (SEQ ID NO:1)	108	120	106	108	n.s.

^a mean ± SEM of at least three determinations

^b Peptide 1 caused significant inhibition only when added to the lower compartment
n.s. = no statistically significant inhibition (p > 0.05)

Furthermore, peptide 3(1-12)[MCP-1] having SEQ ID NO:1 (as well as peptide 1[MCP-1] (SEQ ID NO:2) and peptide 2(1-15)[MCP-1] (SEQ ID NO:3)) did not significantly inhibit THP-1 migration induced by TGF-beta even at 100 μ M. Taken together, these results demonstrated that peptide 3(1-12)[MCP-1] (SEQ ID NO:1) is a general and specific inhibitor of chemokine signaling. Although peptide 3(1-12)[MCP-1] (SEQ ID NO:1) shows weak selectivity for CC chemokines over CXC chemokines, nevertheless, at 100 μ M, peptide 3(1-12)[MCP-1] (SEQ ID NO:1) inhibits >99% of the migration induced by any of the chemokines of either chemokine family tested (Table 2). Thus, although MCP-1 signals through multiple related receptors, peptide 3(1-12)[MCP-1] (SEQ ID NO:1) blocks all of the receptors which participate in the chemotactic and mitogenic signaling pathways elicited by MCP-1.

To exclude the possibility that peptide 3(1-12)[MCP-1] (SEQ ID NO:1) was more effective on THP-1 cells than primary human monocytes, the effect of peptide 3(1-12)[MCP-1] (SEQ ID NO:1) on the chemokine-induced migration of freshly prepared peripheral blood monocytes from 3 donors was tested. Similar to the results for THP-1 cells, 100 μ M of peptide 3(1-12)[MCP-1] (SEQ ID NO:1), but not peptide 1[MCP-1] (SEQ ID NO:2) or peptide 2(1-15)[MCP-1] (SEQ ID NO:3), inhibited all or almost all (>95%) of the migration induced with each of the four chemokines, but did not affect TGF-beta induced migration. Thus, peptide 3(1-12)[MCP-1] (SEQ ID NO:1) is an inhibitor of a broad range of pro-inflammatory chemokines which act on a wide range of target cells (smooth muscle cells, THP-1, Jurkat T-cell line and primary human monocytes). Note that in contrast to THP-1 cells, peptide 2(1-15)[MCP-1] (SEQ ID NO:3) inhibition of MCP-1 induced migration of primary human monocytes (20%) was statistically significant (Table 2).

Example 2

Characterization of Fragments and Variants of Peptide 3(1-12)[MCP-1]
and Peptide 2 [MCP-1]

To determine whether a fragment of peptide 3 has biological activity and 5 selectivity, two 6mer “half-peptides” were analyzed (Table 3): EICADP (SEQ ID NO:8), corresponding to peptide 3(1-6)[MCP-1], and KQKWVQ (SEQ ID NO:9), corresponding to peptide 3(7-12)[MCP-1]. Peptide 3(7-12)[MCP-1] (SEQ ID NO:8) was as potent an inhibitor of CC chemokine signaling as peptide 3(1-12)[MCP-1] (SEQ ID NO:1), but was noticeably more potent as an inhibitor of CXC 10 chemokines (Table 4). In contrast, peptide 3(1-6)[MCP-1] (SEQ ID NO:8) was much less potent as an inhibitor than peptide 3(1-12)[MCP-1] (SEQ ID NO:1).

TABLE 3

	NAME	SEQUENCE	SOURCE
15	<u>Peptide 1 family</u>		
	Pep1	AQPDAINAPVTCC (SEQ ID NO:2)	Residues 1-13 of mature hMCP-1
	<u>Peptide 2 family</u>		
	Pep2(1-15)[MCP1]	SYRRITSSKCPKEAV (SEQ ID NO:3)	Residues 28-42 of mature hMCP-1
	Pep2(1-15)[SDF1]	HLKILNTPNCALQIV (SEQ ID NO:4)	Residues 26-40 of mature hSDF-1 β
20	Pep2(1-14)[MIP1 α]	DYFETSSQCSKPGV (SEQ ID NO:5)	Residues 28-41 of mature hMIP1 α
	Pep2(1-16)[IL8]	ELRVIESGPHCANTEI (SEQ ID NO:6)	Residues 27-42 of mature hIL-8
	<u>Peptide 3 family</u>		
	Pep3(1-12)[MCP-1]	EICADPKQKWVQ (SEQ ID NO:1)	Residues 50-61 of mature hMCP-1
	Pep3(3-12)[MCP-1]	CADPKQKWVQ (SEQ ID NO:7)	Residues 52-61 of mature hMCP-1
25	Pep3(1-6)[MCP-1]	EICADP (SEQ ID NO:8)	Residues 50-55 of mature hMCP-1
	Pep3(7-12)[MCP-1]	KQKWVQ (SEQ ID NO:9)	Residues 56-61 of mature hMCP-1

NAME	SEQUENCE	SOURCE
Leu ₄ Pep3 (1-12)[MCP-1]	EICLDPKQKWVQ (SEQ ID NO:10)	Mutant of peptide 3
Ser ₇ Pep3 (1-12)[MCP-1]	EICADPSQKWVQ (SEQ ID NO:11)	Mutant of peptide 3
5 Ser ₇ Glu ₈ Glu ₉ Pep3 (1-12)[MCP-1]	EICADPSEEWVQ (SEQ ID NO:12)	Residues 50-61 of mature hMIP1 α
Ile ₁₁ Pep3 (1-12)[MCP-1]	EICADPKQKWIQ (SEQ ID NO:13)	Mutant of peptide 3
10 Leu ₄ Ile ₁₁ Pep3 (1-12)[MCP-1]	EICLDPKQKWIQ (SEQ ID NO:14)	Mutant of peptide 3
<u>Unrelated control peptide</u>		
Peptide C	CPSLEDSFIQVA (SEQ ID NO:15)	C-terminus of h Apo(a)RG-C protein
15 D-ala-peptide 3(1-12)[MCP-1]		

20 TABLE 4
Effect of Mutant Sequence Peptide 3 Derivatives on THP-1 Migration

PEPTIDE	ED ₅₀ (μ M) versus				
	MCP1	MIP1 α	IL8	SDF1 α	TGF β 1
Peptide 3 (SEQ ID NO:1)	8	8	14	10	n.s.
Peptide 3[3-12] (SEQ ID NO:7)	8	7	9	9	n.s.
25 Peptide 3[1-6] (SEQ ID NO:8)	33	25	17	19	n.s.
Peptide 3[7-12] (SEQ ID NO:9)	7	5	6	6	n.s.
Leu ₄ peptide 3 (SEQ ID NO:10)	8	7	3	3	n.s.
Ser ₇ peptide 3 (SEQ ID NO:11)	7	6	3	4	n.s.
Ile ₁₁ peptide 3 (SEQ ID NO:13)	6	4	2	7	n.s.
30 Leu ₄ Ile ₁₁ peptide 3 (SEQ ID NO:14)	2	1	3	3	n.s.
Ser ₇ Glu ₈ Glu ₉ pep3 (SEQ ID NO:12)	7	2	9	5	n.s.
WVQ	8	<1	<1	<1	n.s.
KQK	7	n.s.	n.s.	n.s.	n.s.
SEE	n.s.	6	n.s.	n.s.	n.s.

Peptide 3(7-12)[MCP-1] (SEQ ID NO:9) showed essentially no selectivity, inhibiting migration by all chemokines tested with an ED₅₀ in the range of 7-9 µM. Peptide 3(1-6)[MCP-1] (SEQ ID NO:8) was much less efficient at inhibiting the CC chemokines (ED₅₀ of about 30 µM) but only slightly less efficient at inhibiting CXC chemokines (18 µM) compared with peptide 3(1-12)[MCP-1] (SEQ ID NO:1). The selectivity ratio is defined as the average ED₅₀ for MCP-1 and MIP1 α divided by the average ED₅₀ for IL-8 and SDF1 α . Selectivity ratios of greater than 1 indicate greater inhibition of CC chemokines relative to CXC chemokines; selectivity ratios of less than 1 indicate greater inhibition of CXC chemokines relative to CC chemokines; and a selectivity ratio of 1 indicates that both families of cytokines are inhibited to the same extent. Hence, although it is overall a markedly weaker inhibitor of chemokine signaling, peptide 3(1-6)[MCP-1] (SEQ ID NO:8) showed a 2-fold selectivity for CXC chemokines. Thus, peptide 3(1-6)[MCP-1] (SEQ ID NO:8) is a preferred inhibitor of the CXC chemokines, with a selectivity ratio of 0.7, while peptide 3(7-12)[MCP-1] (SEQ ID NO:9) is a preferred inhibitor of both classes of chemokines, with a selectivity ratio of 1.1. The selectivity ratio for peptide 3(1-12)[MCP-1] (SEQ ID NO:1) is 1.5.

Peptide 3(3-12)[MCP-1] (SEQ ID NO:7) had very similar properties to peptide 3(1-12)[MCP-1] (SEQ ID NO:1). This result suggested that the glutamate (E) and isoleucine (I) residues at positions 1 and 2 of the peptide 3(1-12)[MCP-1] (SEQ ID NO:1) sequence, which are not conserved in chemokine sequences other than MCP-1, are unimportant for receptor binding. Alignment of all human chemokine sequences in the peptide 3 region indicate a common conserved motif present in almost all chemokines whether of the alpha or beta subfamily (Table 3).
25 This motif is: Cx₁DPx₂x₃x₄Wx₅Q.

Furthermore, there is a pattern of amino acids in the variable positions x₁ through x₅, which suggests that the nature of the amino acid at these positions may play a role in determining the selectivity of receptor binding. For example, in the CC chemokine family, position x₁ is usually occupied by alanine (A), whereas this

position in commonly leucine (L) in the CXC chemokines except in SDF1 (Isoleucine (I) in SDF-1). To test this hypothesis, the selectivity of Leu₄peptide 3(1-12)[MCP-1] (SEQ ID NO:10) was compared to peptide 3(1-12)[MCP-1] (SEQ ID NO:1). While Leu₄peptide 3(1-12)[MCP-1] (SEQ ID NO:10) showed an 5 approximately 4-fold increase in potency as an inhibitor of CXC chemokines compared with ala-containing peptide 3(1-12)[MCP-1] (SEQ ID NO:1), there was no decrease in the potency of CC chemokine inhibition (Table 4). Thus, Leu₄peptide 3(1-12)[MCP-1] (SEQ ID NO:10) showed some CXC selectivity (a selectivity ratio of 0.37) and was the most CXC selective of all the derivatives tested 10 other than the tripeptides (see below).

As noted for position x₁ above, only three different amino acids appear at position x₅ (Table 1). Most chemokines have valine (V) at position x₅, as do the CXC chemokines IL-8 and MIP. In contrast, SDF-1 and IP10 have isoleucine (I) at this position, while ENA78 is the only chemokine with leucine (L) at this position. 15 The results showed that Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:13) showed some CXC selectivity, though not as marked as Leu₄peptide 3(1-12)[MCP-1] (SEQ ID NO:10) (a selectivity ratio of 0.9), but surprisingly showed the greatest selectivity for IL-8 (which has valine at this position) not SDF-1. This analog was the most selective inhibitor of IL-8 signaling other than the tripeptides, i.e., the analog had a 20 selectivity of IL-8 over other chemokines by about 3 fold.

An analog having both the Leu₄ and Ile₁₁ substitutions did not show any greater specificity as an inhibitor of CXC chemokines than either single mutant Leu₄peptide 3(1-12)[MCP-1] (SEQ ID NO:10) or Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:13) (Table 6). However, Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14) was approximately 5-fold more potent as an inhibitor of CC chemokines 25 than peptide 3(1-12)[MCP-1] (SEQ ID NO:1), or the single mutants Leu₄peptide 3(1-12)[MCP-1] (SEQ ID NO:10) or Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:13). Thus, Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14) was a more potent general chemokine inhibitor, with an average ED₅₀ of 2.3 μM compared with 10 μM for

peptide 3(1-12)[MCP-1] (SEQ ID NO:1). Furthermore, the Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14) unexpectedly preserved the modest CC selectivity of peptide 3(1-12)[MCP-1] (SEQ ID NO:1) with a selectivity ratio of 2.0. Surprisingly, therefore, the Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:1) was 5 approximately 5-fold more potent as an inhibitor of MCP-1 signaling than peptide 3(1-12)[MCP-1] (SEQ ID NO:1), despite the fact that peptide 3(1-12)[MCP-1] (SEQ ID NO:1) contains the cognate sequence from human MCP-1. Moreover, it was found that the Leu₄Ile₁₁peptide 3(3-12)[MCP-1] (SEQ ID NO:1), like 10 Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14), was a higher affinity peptide analog of peptide 3(1-12)[MCP-1] (SEQ ID NO:1).

For positions x₂ through x₄, all chemokines described to date have at least one charged amino acid in this tripeptide region (Table 1). Many chemokines have two basic residues occupying x₂ and x₄ (e.g., KQK in MCP-1, KER in MCP-2 and KLK in SDF-1) while others have two acidic residues (e.g., SEE in MIP1 α , SES in 15 MIP1 β , and SES in RANTES). A recent report (*Nature Med.*, 3, 367 (1997)) suggested that the charge in the extracellular loops of the chemokine receptors may be an important determinant of ligand specificity, e.g., CXCR4 which binds SDF-1 is negatively charged, while CCR5, which binds MIP1 α , MIP1 β and RANTES is positively charged. Thus, residues x₂-x₄ may play an important role in receptor 20 specificity.

To test this hypothesis, several variants were prepared: Ser₇peptide 3(1-12)[MCP-1] (SEQ ID NO:11) substitutes the positively charged K residue present in MCP-1, MCP-2, Eotaxin, IL-8 and SDF-1 with the hydroxylated S residue present in MIP1 α , MIP1 β and RANTES. However, this alteration did not markedly alter 25 the selectivity. In particular, this alteration did not decrease the potency of inhibition of MCP-1 signaling, nor increase the potency of inhibition of MIP1 α signaling (Table 4). The only detectable change was a modest shift from the moderate CC selectivity of peptide 3(1-12)[MCP-1] (SEQ ID NO:1) to a moderate CXC selectivity of the Ser₇peptide 3(1-12)[MCP-1] (SEQ ID NO:11) variant (a

selectivity ratio of 0.5). Another variant, Ser₇Glu₈Glu₉peptide 3(1-12)[MCP-1] (SEQ ID NO:12) which converts the peptide from being the cognate of the MCP-1 sequence to the cognate of the MIP1 α sequence, resulted in a more selective MIP1 α inhibitor, although the selectivity ratio for MIP1 α versus all other chemokines was 5 only about 3 fold.

None of the peptide 3(1-12)[MCP-1] variants had any detectable activity as an inhibitor of TGF-beta induced migration of THP-1 cells, even at 100 μ M (Table 4). Thus, all these variants were highly selective inhibitors of chemokine-induced signaling. There were no substitutions which altered an amino acid residue 10 in peptide 3(1-12)[MCP-1] to any other amino acid regions found in the chemokine sequences described above which markedly reduced the potency of the general chemokine inhibition observed. However, certain alterations resulted in a marked shift in selectivity. For example, the CC selectivity of peptide 3(1-12)[MCP-1] (SEQ ID NO:1) can be converted to CXC selectivity by mutating A to L at position 15 4 (x_1) or by mutating V to I at position 11 (x_5). In particular, two variants had greater than 3-fold selectivity for one chemokine over the average ED₅₀ for all the others, i.e., Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:13) had weak overall selectivity for IL8 inhibition and Ser₇Glu₈Glu₉peptide 3(1-12)[MCP-1] (SEQ ID NO:12) had weak overall selectivity for MIP1 α .

20 In summary, although peptide 3(1-12)[MCP-1] variants varied to a small extent in their ED₅₀s and their specificity for either the α family or β family of chemokines, nevertheless, they were all similar to peptide 3(1-12)[MCP-1] (SEQ ID NO:1). The results in Table 6 showed that peptide 3(1-12)[MCP-1] (SEQ ID NO:1) and peptide 3(1-12)[MCP-1] variants inhibited migration induced by MCP-1, 25 MIP1 α , IL8 and SDF1 α chemokines to a similar extent. While some peptides or peptide variants showed slight preference for CC chemokines, others showed slight preference for CXC chemokines but in no case did the CC-specificity exceed two-fold. Peptide 3(1-12)[MCP-1] (SEQ ID NO:1), peptide 3(1-6)[MCP-1] (SEQ ID

NO:8) and peptide 3(7-12)[MCP-1] (SEQ ID NO:9) also showed no significant CC or CXC selectivity.

Example 3

5 Identification and Characterization of Chemokine Sequences

that are Functional Agonists

The results described above, that peptide 2(1-15)[MCP-1] (SEQ ID NO:3) did not inhibit MCP-1 activity in the THP-1 migration assay, did not exclude the possibility that peptide 2(1-15)[MCP-1] (SEQ ID NO:3) associates with the MCP-1 receptor and either fails to inhibit MCP-1 binding and signaling, or acts as an agonist, preventing binding of MCP-1 but transducing an MCP-1 like signal. To determine whether peptide 2(1-15)[MCP-1] (SEQ ID NO:3) binds to chemokine receptors, THP-1 cells were mixed with a biotinylated derivative of peptide. Peptide 2(1-15)[MCP-1] (SEQ ID NO:3) was found to bind to THP-1 cells with a reasonable affinity ($kD = 1.9 \mu M$), suggesting that peptide 2(1-15)[MCP-1] (SEQ ID NO:3) was able to interact with chemokine receptors without inhibiting chemokine signaling (a neutral binding agent).

Unlike peptide 3, which represents a region that is relatively conserved between chemokines, there is much less marked sequence similarity in the peptide 2 region of chemokines. Thus, peptide 2, derivatives or variants thereof, may possess more chemokine-specific effects. To test this hypothesis, the binding of biotinylated peptide 2(1-15)[MCP-1] (SEQ ID NO:3) to two different cell types, i.e., THP-1 cells and Jurkat cells, was compared. THP-1 cells express receptors for MCP-1, MIP1 α , SDF-1 α and IL-8 while Jurkat cells express functional receptors for SDF-1 only. Peptide 2(1-15)[MCP-1] (SEQ ID NO:3) bound to Jurkat cells with a similar kD ($3 \mu M$) to THP-1 cells ($1.9 \mu M$). This observation suggests that, surprisingly, peptide 2(1-15)[MCP-1] was able to bind to a number of different chemokine receptors, although the sequence of peptide 2(1-15)[MCP-1] (SEQ ID NO:3) shows little homology to the equivalent region from SDF-1.

The functional agonist properties of peptide 2(1-15)[MCP-1] (SEQ ID NO:3) were characterized by incubating THP-1 cells with varying concentrations of peptide 2(1-15)[MCP-1] (SEQ ID NO:3) in a migration assay. Peptide 2(1-15)[MCP-1] (SEQ ID NO:3) had weak agonist activity (promoting migration with an ED_{50} of about 10 μ M) with maximal migration at 100 μ M, which was approximately 10% of that induced by MCP-1. Thus, at high concentrations (>20 μ M) peptide 2(1-15)[MCP-1] (SEQ ID NO:3), variants or derivatives thereof, may be useful for applications which require weak MCP-1 agonist activity (for example, for the therapy of parasitic infections where increased macrophage activity is desirable). Moreover, at lower concentrations (> 1 μ M but < 20 μ M), peptide 2, peptide 2 variants or derivatives may be useful as a neutral binding agent which may affect binding of non-MCP-1 proteins to the chemokine receptors. For example, peptide 2, derivatives or variants thereof, may be useful to prevent or inhibit HIV binding to chemokine receptors without inhibiting desirable chemokine signaling.

15

Example 4

Identification, Preparation and Characterization of Therapeutic Agents of the Invention for *In Vivo* Use

A. Derivatives

20 Peptides are generally susceptible to chemical or enzymatic hydrolysis. In particular, peptides are not normally bioavailable by the oral route since they are not stable in the acid and proteolytic environment of the stomach. Thus, chemical or enzymatic hydrolysis leads to a very short *in vivo* half-life for peptides. To extend the half-life of agents susceptible to hydrolysis, *in vitro* active agents are modified in 25 a manner that results in a derivative which may be orally bioavailable, have improved pharmacokinetics, and the administration of which may achieve concentrations in blood that inhibit chemokine activity. For example, cyclic-reverse-D (CRD) peptides may be prepared. CRD peptides are prepared by synthesizing the reverse sequence of the peptide (C-terminal to N-terminal) using

the opposite stereoisomer (D-amino acids in place of L amino acids). The resulting peptide is then cyclized via N- and C-terminal cysteine residues. These derivatives retain a very similar steric arrangement of atoms to non-CRD peptide, but are not subject to enzymatic hydrolysis. Other derivatives which may exhibit an extended 5 half-life *in vivo* include thienyl or pyridyl derivatives (e.g., U.S. Patent No. 4,992,463; U.S. Patent No. 5,091,396).

For example, to prepare a peptide 3 derivative, peptide 3(3-12)[MCP-1] was modified according to Jameson et al. (*Nature*, **368**, 744 (1994)), which yielded CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1] (Figure 4). CRD-Cys₁₃Leu₄Ile₁₁peptide 10 3(3-12)[MCP-1], which had very similar properties to peptide 3(1-12)[MCP-1] (SEQ ID NO:1) in the *in vitro* assays described hereinabove, was found to be stable against both acid hydrolysis (<10% degradation at pH 2.0 for 2 hrs at 37°C) and enzymatic destruction (5 units trypsin for 2 hrs at 37°C). CRD-Cys₁₃Leu₄Ile₁₁peptide 15 3(3-12)[MCP-1] was also resistant to hydrolysis *in vivo* and allowed therapeutically useful plasma concentrations to be achieved (> 10 μM 24 hours after a single intraperitoneal dose of 1 mg of CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1] in 250 μl saline).

Cyclic-reverse D (CRD), linear reverse-D (LRD), cyclic forward L (CFL), and linear forward L (LFL) (i.e., the standard form of peptides) derivatives of 20 Leu₄Ile₁₁peptide 3 were prepared and their MCP-1 inhibitory activity in the THP-1 transwell assay determined. The results were

	LFL-Leu ₄ Ile ₁₁ peptide 3	1-5 μM
	LRD-Leu ₄ Ile ₁₁ peptide 3	200-400 nM
25	CFL-Cys ₁₃ Leu ₄ Ile ₁₁ peptide 3	500-700 nM
	CRD-Cys ₁₃ Leu ₄ Ile ₁₁ peptide 3	50-100 nM

These results show, somewhat surprisingly, that both cyclization and reverse-D derivatization independently improve activity. This improvement is then

additive in the CRD derivative. Thus, cyclization improved affinity by constraining the conformations of the peptide. However, it was not expected that the reverse-D derivatization would be so beneficial, possibly by increasing stability of the molecule.

5 CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1] was found to be a very potent inhibitor of MCP-1 induced THP-1 migration (ED₅₀ of about 1 nM). This increased potency compared to the parent Leu₄Ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14) may reflect increased stability, even *in vitro*, or it may reflect the increased conformational stability of the peptide. Moreover, this compound binds to the 10 signaling receptor with the same affinity as native full-length MCP-1 but does not signal.

To determine if CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] inhibited or enhanced the proliferation of T or B cells to conconavalin A or tetanus toxoid in culture, proliferation of CD4 T cells and B cells was assessed by CFSE-FITC cell 15 labeling. 50 ng of CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12) [MCP-1] inhibited ConA proliferation of CD4 T cells by 50% and 5 ng of CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12) [MCP-1] reduced ConA proliferation of CD4 T cells by < 3%. CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] had no effect on proliferation of B cells to tetanus toxoid.

20 Computer modeling was employed to determine whether specific amino acid replacements affected the conformation of the peptide derivative CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1]. The peptide sequence was entered into HyperChem 5.0 (HyperCube). A minimum energy conformation was sought using the Amber Force Field parameters and the Polak-Ribiere algorithm. The initial 25 model was manipulated both by molecular dynamics simulations (300°K, 2 nsec) and manual sidechain rotations, followed by geometry optimization, until an apparent global minimum energy conformation was reached. Convergence criterion was <0.01 Kcal/mol Å. A conformation was obtained using this procedure with an energy of about 213.4 kcal/mol.

To test the sensitivity of the model peptide to perturbations, each of the residues except the terminal cysteines forming the disulfide bond was mutated individually from D to L, and the geometry re-optimized, starting with the minimum conformation of the all D peptide. For these perturbations each mutant was first run 5 through the geometry optimization routine, then a molecular dynamics simulation, then another geometry optimization. The resulting mutant peptides were compared to the all-D form by overlaying the disulfide bond and one adjacent atom, and visually assessing the difference between the peptide backbones. The overall conformation was insensitive to change of chirality at positions 2, 3, 4, 8, 9, and 10, 10 but was sensitive to change of chirality at positions 5, 6, and 7. Generally, changes in sidechain position were minor except when the backbone conformation changed significantly. Energies for the mutants varied from -187.9 to -226.1 Kcal/mol, but the energy change (from -213.4 for the starting conformation) did not correlate with conformational change.

15 In addition, the effect of modifying the aspartate residue at position 9 was examined by converting its sidechain carboxyl group to the D-alanyl amide. A minimum energy conformation of the modified peptide was sought using the same routine as for the chiral mutants, starting from the same minimum energy conformation. Condensation of D-alanine to the residue 9 sidechain carboxyl 20 caused a major change in the conformation of the peptide. This is consistent with the *in vitro* monocyte migration data which demonstrated a significant loss in biological activity of the D-ala peptide relative to CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1].

Molecular modeling indicated that L-Leu-CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-25 12)[MCP-1], which is CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] with the D-Leu replaced with an L-Leu, should result in very little change to the conformation of the peptide backbone. *In vitro* migration studies with the L-Leu-derivative showed that it retained functional activity as well. Thus, to select for particular amino acid

substitutions which retain the conformation of a biologically active molecule of the invention, molecular modeling may be employed.

The following D-amino acid to L-amino acid changes had no significant impact on the structure of the peptide backbone as assessed by modeling.

	<u>Amino Acid</u>	<u>Position</u>	<u>kcal/mole</u>
5	GLN	2	-205.5
	ILE	3	-202.4
	TRP	4	-222.0
	PRO	8	-226.1
10	LEU	10	-211.9

The following D-amino acid to L-amino acid changes had a significant impact on the structure of the peptide backbone as assessed by this technique.

	<u>Amino Acid</u>	<u>Position</u>	<u>kcal/mole</u>
15	LYS	5	-200.1
	GLN	6	-211.6
	LYS	7	-187.9
	ASP	9	-214.9

20 B. DARC binding

A further consideration for bioavailability is non-specific binding of the therapeutic agent. Red blood cells have a signaling-deficient chemokine receptor or binding protein, termed the Duffy Antigen Receptor for Chemokines (DARC).

Although it does not signal, this receptor has a high affinity for chemokines (10 nM)

25 and may play a role in clearing them from the circulation. Unfortunately, any chemokine receptor antagonist which has a high affinity for DARC may be sequestered by the huge pool of binding sites on red blood cells, and hence unavailable to inhibit productive chemokine signaling in other tissues. Similarly, agonists which bind DARC with high affinity are unavailable to productively signal
30 through specific chemokine receptors. For *in vivo* use, an agent of the invention preferably has some affinity for DARC, since peptides which do not bind to DARC

are rapidly cleared at first pass by glomerular filtration. Thus, preferred agents have DARC binding (affinity constant) in the range 100 nM to 1 mM, more preferably in the range 1 μ M to 100 μ M and even more preferably in the range of 10 to 100 μ M.

Although the interaction of chemokines with DARC is high affinity
5 (5-10 nM association constant), kinetically the interaction is characterized by extremely rapid on and off rates. Consequently, incubation with labeled chemokine leads to saturation of the DARC binding sites, but most of the bound label is lost within minutes of removing the unbound label (>90% loss within 3 minutes). As a result, it is difficult to directly determine the binding of peptides to DARC by
10 assaying direct binding of biotinylated peptide, since the rapid off rates make determination of the amount of bound label impossible or inaccurate.

To overcome this difficulty, the k_a for association of DARC with peptide
3(1-12)[MCP-1] (SEQ ID NO:1) and peptide 2(1-15)[MCP-1] (SEQ ID NO:3) was estimated by incubating red blood cells expressing DARC with 125 I-labeled MCP-1
15 in the presence of varying concentrations of peptide. After binding has reached equilibrium (30 minutes at 37°C), the cells are separated from the unbound label by centrifugation for 5 minutes through a sucrose gradient. Counts associated with the cells are then determined by gamma-counting scintigraphy. In the absence of all peptides, the association constant for 125 I-labeled MCP-1 on human red blood cells
20 was 5.45 nM, a value which is in accord with a previous report. Furthermore, Scatchard analysis confirms the presence of a single high affinity binding site with 500-1000 copies per cell, consistent with the known properties of DARC. Thus, determination of 125 I-MCP-1 binding to red blood cells in this assay in the presence of various concentrations of the peptide(s) allows the association constant of the
25 peptide for DARC to be accurately estimated.

The DARC specificity ratio is also determined. The DARC specificity ratio is defined as the estimated k_a for association with DARC divided by the ED_{50} for biological activity. A DARC specificity ratio greater than 1 indicates that a peptide associates poorly with DARC and is bioavailable for modulating chemokine

signaling, either as an antagonist or agonist. A DARC specificity ratio of about 1 indicates that the peptide binds DARC and the THP-1 signaling receptors with similar affinity. Thus, it may be difficult to achieve biologically active (as a chemokine inhibitor) concentrations of these peptides *in vivo* without further 5 modifications of the peptide. A DARC specificity ratio less than 1 indicates much higher affinity for DARC than for chemokine signaling receptors.

Peptide 1[MCP-1] (SEQ ID NO:2)(which does not bind to chemokine receptors but functions in a dominant negative fashion) showed no binding to DARC (estimated $k_a > 100 \mu\text{M}$). In marked contrast, the weak agonist peptide 2(1-10 15)[MCP-1] (SEQ ID NO:3) showed high affinity binding to DARC. The association constant for peptide 2(1-15)[MCP1] (SEQ ID NO:3) for chemokine receptors on THP-1 cells was estimated at 2 μM using competition binding analysis. However, this peptide had an affinity for DARC of less than 500 nM, also assessed by competition binding analysis, using red blood cells. Thus, peptide 2(1-15)[MCP1] (SEQ ID NO:3) binds to THP-1 cell chemokine receptors, although it does not inhibit signaling through the receptors, and it binds DARC even more strongly (DARC selectivity ratio = 0.1-0.2). Thus, peptide 2 is a preferred therapeutic agent for the treatment or prevention of malaria (an action requiring DARC inhibition, but not modulation of chemokine signaling).

Peptides, such as peptide 2(1-15)[MCP-1] (SEQ ID NO:3) which have very high affinity for the DARC receptor, may have strong biological agonist activity *in vivo* (although they are only weak agonists or neutral agonists *in vitro*). Moreover, peptide 2, variants and derivatives thereof may be strongly pro-inflammatory *in vivo*, or strongly exacerbate existing inflammation by preventing DARC from performing 20 the function of binding chemokines. If DARC functions as a sink to remove chemokines from the circulation, then the concentration of chemokines may be markedly increased by the presence of peptide 2. Under conditions where 25 chemokines are being made a released into the circulation (e.g., during inflammation), peptide 2 may exacerbate that inflammation, allow the inflammation

to persist longer than in the absence of the peptide or otherwise change the qualitative nature of the inflammatory reaction. For these reasons, peptides with a low DARC specificity ratio are useful for the treatment of conditions which require improved immune function, or conditions which are characterized by a 5 pathologically inadequate inflammatory response.

MIP1- α has previously been shown to be the only chemokine which does not bind with significant affinity to DARC. Peptide 2(1-9)[MCP-1] had a Duffy affinity of about 50 μ M while peptide 2(1-15)[MIP1- α] (SEQ ID NO:5) was a potent receptor binding agent for the MIP1- α receptor and had excellent specificity over 10 DARC. That is, peptide 2(1-15)[MIP1 α] (SEQ ID NO:5) did not bind to DARC (association constant > 50 μ M) but bound strongly to chemokine receptors on THP-1 cells (association constant = 100-900 nM; number of binding sites is about 150,000 per cell). Moreover, this agent did not inhibit THP-1 cell migration 15 induced by MCP-1, MIP1 α , IL-8, or SDF1 α . Thus, this latter agent may be particularly useful as a neutral chemokine receptor binding agent *in vivo*, highly selective over DARC.

Peptide 3(1-12)[MCP-1] (SEQ ID NO:1) also binds to DARC, although it binds to DARC with only a similar affinity to which it binds to the chemokine receptors (low μ M concentration range). Leu₄ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID 20 NO:14) had essentially no DARC binding capacity, while inhibiting MCP1 induced migration at concentrations around 1 μ M. Thus, peptide 3 derivatives, such as leu₄ile₁₁peptide 3(1-12)[MCP-1] (SEQ ID NO:14) may achieve antagonist properties *in vivo*.

The shorter fragments of peptide 3[MCP-1] (e.g., peptide 3(7-12)[MCP-1] 25 (SEQ ID NO:9)) showed progressively higher DARC specificity ratios (about 3.0 for peptide 3(7-12)[MCP-1] (SEQ ID NO:9) versus 1.0 for peptide 3(1-12)[MCP-1] (SEQ ID NO:1)), indicating that where chemokine signaling receptor specificity is desired, shorter peptide fragments which retain full chemokine antagonist or agonist activity are in general to be preferred over the full length peptides.

Peptide 3(1-12)[MCP-1] (SEQ ID NO:1) (DARC specificity ratio = 1.00) is unlikely to be useful as a pan-chemokine inhibitor *in vivo*, whereas the Leu₄Ile₁₁peptide 3[MCP-1] (SEQ ID NO:14) (DARC specificity ratio = 37.83), or its derivatives such as CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1], which bound only

5 weakly to DARC (association constant = 90 μ M) but bound very strongly to chemokine receptors on THP-1 cells (association constant = 100-500 nM; number of binding sites is about 150,000 per cell), are a preferred embodiment for the treatment or prevention of atherosclerosis, osteoporosis, and autoimmune diseases, and HIV infection (chemokine signaling receptor binding functions). Moreover,

10 CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1] inhibited THP-1 cell migration induced by MCP-1, MIP α , IL-8, and SDF α , with very similar ED₅₀s.

CRD-peptide 2(1-15)[MCP-1] has more functional potency, less Duffy binding activity compared with the LFL derivative. LRD peptide 2(1-15)[MCP-1] had approximately a 100-fold decrease in Duffy binding (25 μ M versus 100 μ M for

15 LFL).

An alternative approach to preparing agents that are bioavailable is the preparation of non-peptide analogs of chemokines. A preferred non-peptide analog of the invention includes an isostere of WIQ, e.g., a compound of formula (IV), wherein Z=CH₃; Y=O; X=CH₃; and Ar = indolyl. This compound did not bind to

20 DARC (association constant = > 30 μ M) but bound very strongly to chemokine receptors to THP-1 cells (association constant = 100 nM-1 μ M; number of binding sites is about 150,000 per cell). This agent inhibited THP-1 cell migration induced by MCP-1, MIP α , IL-8 and SDF α with very similar ED₅₀s.

TABLE 5

PEPTIDE	SEQUENCE	SPECIFICITY*	DUFFY SELECTIVITY ^b	AVERAGE ED ₅₀ (μM) ^c
Pep2[MCP1]	SYRRITSSKCPKEAV (SEQ ID NO:3)	-	0.18 ^d	2 ^e
Pep2[SDFF1]	HLKILNTPNCALQIV (SEQ ID NO:4)	-	<1 ^d	1-10 ^e
Pep2[MIP1 α]	DYFETSSQCSPKPGV (SEQ ID NO:5)	-	>100 ^d	1-10 ^e
Pep2[IL8]	ELRVIESGPHCANTEI (SEQ ID NO:6)	-	<1 ^d	1-10 ^e
Pep3	EICADPKQKWWVQ (SEQ ID NO:1)	1.5	1.00	10
Pep3[3-12]	CADPKQKWWVQ (SEQ ID NO:7)	1.2	1.21	8
Pep3[1-6]	EICADP (SEQ ID NO:8)	0.7	5.32	24
Pep3[7-12]	KQKWWVQ (SEQ ID NO:9)	1.0	2.94	6
Leu ₄ pep3	EICLDPKQKWWVQ (SEQ ID NO:10)	0.4	n.d.	5
Ser ₇ pep3	EICADPSQKWWVQ (SEQ ID NO:11)	0.5	2.00	5
Ile ₁₁ pep3	EICADPKQKWIQ (SEQ ID NO:12)	0.9	n.d.	5
Leu ₄ Ile ₁₁ pep3	EICLDPKQKWIQ (SEQ ID NO:13)	2.0	37.83	2
Ser ₇ Glu ₈ Glu ₉	EICADPSEEQVQQ (SEQ ID NO:2)	1.6	0.59	6
-	KQQ (SEQ ID NO:51)	>100	22.34	6 ^f
Pep3[10-12]	WWQ (SEQ ID NO:33)	1.1	24.81	2
-	WIQ (SEQ ID NO:34)	1.0	n.d.	2
-	SEE (SEQ ID NO:27)	>100	n.d.	0.8 ^f
-	KLK (SEQ ID NO:28)	<0.1	n.d.	0.1-10 ^f
-	KEN (SEQ ID NO:55)	<0.1	n.d.	0.1-10 ^f
CRD-Cys ₁₃ pep3[3-12]	-CQIWWKQKPDA--	n.d.	8.82	0.8
LRD-Cys ₁₃ Leu ₄ Ile ₁₁ pep3[3-12]	CQIWWKQKPDA--	n.d.	4.59	1
CRD-Cys ₁₃ Leu ₄ Ile ₁₁ pep3[3-12]	-CQIWWKQKPDA--	n.d.	>100	0.1

Footnotes

- ^a ‘CC-specificity’ is the average inhibitory ED₅₀ versus SDF1 and IL8 divided by average inhibitory ED₅₀ versus MCP-1 and MIP1 α .
- ^b ‘Duffy selectivity’ is the estimated ka for binding to red blood cells divided by average inhibitory ED₅₀ versus each of the chemokines (except for peptide 2; see footnote d below).
- ^c ‘Average ED₅₀’ is the average inhibitory ED₅₀ for inhibition of THP-1 migration induced by each of the chemokines (except for peptide 2; see footnote e below).
- ^d For the peptide 2 family, the ‘average ED₅₀’ is the estimated ka for binding to THP-1 cells.
- ^e For the peptide 2 family, the ‘Duffy selectivity’ is calculated as the ka for binding to red blood cells divided by the ka for binding THP-1 cells.
- ^f For tripeptide derivatives so marked, the peptide is highly specific for one of the four exemplary chemokines. In these cases, the ED₅₀ shown is for inhibition of that chemokine.

n.d. = not determined.

Abbreviations

CRD = Cyclic reverse-D derivative

LRD = Linear reverse-D derivative

CFL = Cyclic derivative of standard L-form peptide

LFL = Standard, linear L-form peptide [NB; all peptides are LFL unless stated otherwise]

Amino acids in italics are D-form amino acids, all others are L-form

-- = Cyclization linking two cysteines so marked

Example 5

Anti-HIV Activity of the Agents of the Invention

To demonstrate that the agents of the invention inhibit HIV binding and infection of cells, human T-cell derived Jurkat cells were incubated with an infectious T-tropic HIV isolate in the presence of (i) no inhibitor, (ii) peptide C (Table 5) as an inactive control peptide, (iii) 100 μ M peptide 3(1-12)[MCP-1] (SEQ ID NO:1), or (iv) 100 ng/ml SDF-1, which should bind to and block all CXCR-4 receptors. After 3 weeks in culture, viral replication was assessed by a reverse transcriptase assay of the culture medium. Peptide 3(1-12)[MCP-1] (SEQ ID NO:1) 10 was found to be an effective inhibitor of Jurkat cell infection by HIV (Figure 3).

Since peptide 2(1-15)[MCP1] (SEQ ID NO:3) binds to chemokine receptors on the surface of Jurkat cells and THP-1 cells, but does not inhibit productive signaling by chemokines, it is possible that peptide 2(1-15)[MCP-1] (SEQ ID NO:3) binds and inhibits an epitope used by HIV for cell entry but not by MCP-1 for 15 signaling. To test this hypothesis, the same HIV infection assay described above was employed to test whether peptide 2(1-15)[MCP-1] (SEQ ID NO:3) inhibits HIV infection of Jurkat cells. At 100 μ M, peptide 2(1-15)[MCP-1] (SEQ ID NO:3) was more effective than peptide 3(1-12)[MCP-1] (SEQ ID NO:1), and as effective as SDF1 α , in preventing virus entry.

20 Peptide 2 derivatives (Figure 10) are better inhibitors of Jurkat T cell infection by HIV (a CXCR4 mediated event) than peptide 3 derivatives, while surprisingly peptide 3 is a better inhibitor of THP-1 cell infection (a CCR-5 mediated event). Thus, combinations of peptide 2 and peptide 3 may be particularly useful for anti-HIV therapy, e.g., to inhibit productive infections by both M-tropic 25 and T-tropic isolates.

Moreover, as LRD peptide 2(1-15)[MCP-1] had a 100 nM affinity constant or lower for CCR5/CXCR4 and a 100 fold decrease in Duffy binding relative to LFL peptide 2[MCP-1], LRD derivatives may be more efficacious than their LFL counterparts (25 μ M versus 100 μ M for LFL).

Current therapies for inhibition of HIV focus on the virus, for example reverse transcriptase inhibitors or viral protease inhibitors. These therapies are only effective for a limited period. In each case, the efficacy is reduced because the virus is undergoing rapid replication, and there is selection in favor of mutants which are 5 resistant to the inhibitors. Although combination therapies are more effective, they are unlikely to result in clearance of the virus from an infected individual. Eventually, mutant virus will arise which circumvents the drug cocktail and progression will again occur in the now drug-resistant individual. Thus, strategies which are based on co-receptor inhibition target a host protein, rather than a virus 10 protein, may have increased efficacy as more extensive mutations in the virus may be necessary to circumvent an inhibited co-receptor. Indeed, the resistance to infection of CCR-5 Δ 32 homozygotes suggests that the virus cannot readily adapt to use of an alternative co-receptor, at least while the virus population is small.

Preferably, a Ser₁₀ variant of peptide 2(1-15)[MCP-1] 15 (SYRRITSSKSPKEAV), or its LRD Cys₆Ser₁₀Cys₁₆ derivative (cvaekpsksstirrysc) or CRD derivative, is employed. DARC binding of SYRRITSSKSPKEAV is in the range 20 μ M to 100 μ M and activity in the range 1-100 nM as an anti-HIV agent.

Example 6

20 Rapid Screening Method for Infectivity

Current assays for HIV infection *in vitro* are time consuming and lack 25 reproducibility. For example, infection is often monitored by the production of viral reverse transcriptase (RT) activity using a radiolabelled RT substrate. Unfortunately, RT production is low, even when a laboratory adapted HIV strain is used to infect a high permissive line such as the Jurkat human T cell line. As a result, it is necessary for the infected cells to be cultured for two or more weeks to allow sufficient infection to occur for RT production be measurable. In addition to being time consuming, this assay has a number of other disadvantages: most importantly, it relies on multiple rounds of secondary infection to increase the viral

titer sufficiently for RT activity to become detectable. As a result, small differences in primary infection are magnified, and since primary infection frequency is low, stochastic differences between identically treated wells become significant. The assay therefore requires many replicate wells for each analysis, with as many as 24 5 replicates being routinely used. For example, in a typical assay groups of 24 wells of Jurkat cells in 96-well plates are infected with replicate aliquots of HIV virus stock, with one group receiving treatment with peptide 2 as a chemokine co-receptor inhibitor, another group receiving SDF-1 α (the CXCR-4 natural ligand) and a third group is untreated. After three weeks, the cells were harvested and RT activity 10 measured. The co-efficient of variation in the untreated wells was 37%. As a result, although peptide 2 inhibited RT activity by 75%, this was significant only with p=0.02 because of the high well to well variability. This necessitates the use of many replicate making the assay cumbersome for screening purposes.

An alternative method is to use direct visualization of the HIV proteins, for 15 example, by immunofluorescence microscopy. Unfortunately, even the most highly expressed HIV proteins (such as p24gag) are present at fairly low levels in cells. Thus, direct detection the earliest stages after infection has been difficult and error prone. Therefore, the following method was employed to enhance the sensitivity of 20 immunofluorescence, allowing the number of HIV infected cells to be accurately determined between 24 hours and 72 hours after infection. Furthermore, the signal to noise ratio of this technique allows automated counting of the infected cells using image analysis software.

For THP-1 cells, the cells are adhered to glass multiwell slides (for example, 16-well chamber slides; Nunc) using PMA and hydrocortisone. The cells are then 25 exposed to virus in the chamber slide in the presence of various test agents. For non-adherent cells such as Jurkat cells, infections are carried out in, for example, 96-well culture plates as for RT assays, but prior to analysis the cells are attached to glass slides using a cyrospin apparatus in accordance with the manufacturer's instructions. The infected cells on the glass slides are fixed between 24 hours and

72 hours after infection, for example, by immersing the slides in ice cold acetone for 90 seconds. Other methods of fixation compatible with quantitative immunofluorescence may also be used (see *J. Histochem. Cytochem.*, **44**, 1043 (1997) for a discussion of quantitative immunofluorescence procedures). Following 5 fixation, non-specific binding of proteins to the cells is blocked, e.g., by incubation in 3% w/v fatty acid free bovine serum albumin in phosphate buffered saline (3% FAF-BSA in PBS) for 30 minutes at room temperature. Alternatively, other blocking solutions (e.g., 5% sucrose, 5% Tween-20 in PBS) may be used. The blocked sections are then stained for HIV protein, for example, using a specific 10 antiserum to p24gag. Slides are incubated with the antiserum at a suitable concentration (usually in the range 1-100 µg/ml of specific IgG) in 3% FAF-BSA in PBS. Antibodies to other HIV antigens may be used, although relatively highly expressed antigens such as p24gag are preferred.

This incubation should be left on for at least 16 hours. Traditional 15 immunofluorescence procedures use primary antibody incubation periods typically 1-2 hours in length, but longer incubation increases signal without increasing background (*J. Histochem. Cytochem.*, **44**, 1043 (1997)). The incubation may be left on for up to 36 hours without deleterious effects on the signal to noise ratio. Unbound antibody is then washed off. Typically, this involves 3 x 3 minute washes 20 in PBS, although other washing regimens may be used (see *J. Histochem. Cytochem.*, **44**, 1043 (1997)) for a comparison of washing methods). Normally, second antibody labelled with an appropriate fluorophore is then used to detect the unbound primary antibody. However, to prevent primary antibody from falling off the antigen, primary antibody is post-fixed to the section. This may be achieved, for 25 example, by incubating the slide in freshly prepared 4% paraformaldehyde in PBS for 10 minutes at room temperature. After three further washes, e.g., 3 x 3 minutes in PBS, the slides are exposed to a secondary antibody specific for the species of the primary antibody coupled to an appropriate fluorophore (for example, antirabbit-IgG FITC conjugate at 1-100 µg/ml). A non-specific nuclear stain should be included in

this incubation. For example, Hoescht 33342 at 1-100 ng/ml could be used, or propidium iodide at 1-100 ng/ml. This incubation is for a minimum of about 4 hours, preferably at least 8 hours and may be left up to 24 hours without detrimental effect on the signal to noise ratio. Slides are then washed, for example, 3 x 5 minutes in PBS, to remove unbound second antibody and mounted with a suitable mounting medium such as Citifluor AF1. Slides are left at least about 18 hours after mounting but less than about 72 hours in a dark box following mounting prior to analysis.

Analysis may be performed manually using any suitable microscope with 10 epifluorescence visualization capability and appropriate filter sets to allow examination of the fluorescence of the secondary antibody fluorophore selected (e.g., FITC) and the non-specific nuclear staining selected (e.g., Hoescht 33342) separately. The number of cells in each field of view is determined by counting nuclei using filters to visualize the non-specific nuclear stain. The number of cells 15 infected with HIV in the same field of view is then determined by switching the filter set to visualize the fluorophore coupled to the secondary antibody. In each case, the number of cells may be determined by manual counting. Alternatively, image analysis software (for example, OpenLab software: Improvision, U.K.) may be used to apply a consistent threshold to each image and count the number of 20 separate objects above that threshold. Deagglomeration algorithms, standard in the field of image analysis, may be applied if required according to the density of the cells on the slides. Provided that constant set of illumination conditions are used during image acquisition and that a constant threshold is applied, the fraction of HIV stained cells may be rapidly and accurately determined without reference to 25 subjective considerations.

Figure 11 shows the results from the inhibition of HIV infection by peptide 2(1-15)[MCP-1] and peptide 3(1-12)[MCP-1]. The coefficient of variation for the technique is less than 5% with excellent reproducibility. Approximately 6% of THP-1 cells were infected by HIV in the absence of peptide inhibitors of the

chemokine receptors. In the presence of 100 μ M peptide 2, this was reduced by 25% \pm 3%. Statistically significant reduction in viral infection can be noted from a single well determination, using multiple fields of view within the single well to establish statistical significance. Positive results can be confirmed by analysis of 5 replicate wells, and dose response curves can be constructed.

Where expression of the HIV antigen selected is low, or where detection very early after infection (from about 14 hours after infection), the sensitivity this technique may be enhanced further. After the secondary antibody incubation, a further post-fixation step is applied (for example, 10 minutes at room temperature in 10 4% paraformaldehyde in PBS) and the slides are incubated with a non-immune immunoglobulin fraction of the same species as the primary antibody (i.e., non-immune rabbit serum). This incubation may be for 1-2 hours at room temperature. Thereafter, a second incubation with the same primary antibody as used previously 15 is then performed. In this case, the non-specific nuclear stain (e.g., Hoescht 33342) is added to the second incubation with secondary antibody only. All incubations should be separated by an appropriate washing regimen (e.g., 3 x 3 minutes in PBS at room temperature). These changes lead to a further five-fold increase in sensitivity for detection of the HIV antigen allowing earlier detection of infection.

20

Example 7

Preparation and Characterization of Tripeptide

Therapeutic Agents of the Invention

To determine whether fragments of peptide 3(1-12)[MCP-1] possessed biological activity, fragments of peptide 3 were prepared. Peptide 3(10-12)[MCP-25 1], i.e., WVQ was found to be a potent inhibitor of all chemokines tested (Table 6). The amino acid residues at positions 10-12 (WVQ) are conserved in many other chemokines, e.g., MCP-3, MIP1 α , MIP1 β , RANTES, EOTAXIN, and IL8, although SDF1 has the sequence WIQ. WVQ inhibited all four of the exemplary chemokines tested, although, unlike peptide 3(1-12)[MCP-1] (SEQ ID NO:1), it was a more

potent inhibitor of all the chemokines other than MCP-1, with ED₅₀s around 1 μ M. Thus, these tripeptides, WVQ and WIQ, as well as non-peptide analogs based on these tripeptides, are pan-specific chemokine inhibitors. Moreover, it was found that WVQ had good Duffy selectivity (i.e., selectivity of 10).

5 Peptide 3(7-9)[MCP-1], i.e., KQK, did not bind to DARC (association constant = > 50 μ M) but bound strongly to chemokine receptors on THP-1 cells (association constant = 500 nM-1 μ M; number of binding sites is about 15,000 per cell). This agent inhibited THP-1 cell migration induced by MCP-1, but did not inhibit migration induced by MIP1 α , IL-8 or SDF1 α . Thus, KQK with an ED₅₀ = 2-5
10 μ M was found to be a specific inhibitor of MCP-1, i.e., it had no effect on MIP1 α , SDF1 α or IL8 induced activity even at 100 μ M. Four tripeptides and a dipeptide of random sequence (RGD, GGR, TTT, APG, and VE) were also tested. None of these significantly inhibited migration induced by any of the chemokines. Thus, the
15 tripeptide KQK was specific for inhibiting MCP-1 activity, showing more than 100-fold specificity for MCP-1 over all the other chemokines tested.

20 Tripeptide equivalents of KQK from MIP1 α , SDF1 α and IL8, based on an alignment of conserved cysteine residues in chemokine sequences, were then tested for their inhibition of chemokine-induced THP-1 migration. In each case, the tripeptide was highly specific for its cognate chemokine (> 100-fold specific in each
25 case). For example, SEE, the cognate peptide from MIP- α , showed greater than 100-fold selectivity for MIP1- α over the other chemokines. Moreover, KLK was a specific and potent inhibitor of SDF1, and KEN was a specific and potent inhibitor of IL8. In no case did the tripeptide significantly inhibit migration induced by any of the non-cognate chemokines, even at 100 μ M. It is envisioned that tripeptides in which a conservative substitution is made may have the same specificity as the native tripeptide. Moreover, the corresponding tripeptides in other chemokines may be specific for their cognate chemokines.

Table 6

Tripeptide	Chemoattractant					
	MCP-1	MIP1 α	RANTES	IL-8	SDF1 α	
5	KQK ^a	95 \pm 8^b	-	-	-	29 \pm 1
	SEE	-	65 \pm 3	-	-	-
	SES	-	-	87 \pm 4	-	-
	KEN	21 \pm 2	-	-	70 \pm 4	-
	KLK	-	-	-		87 \pm 6
10	WVQ ^c	8 μ M	7.5 μ M	1.5 μ M	1 μ M	2 μ M

For each peptide shown (except WVQ), a number indicates the percentage inhibition of migration induced by that chemoattractant by that tripeptide at 100 μ M concentration (mean \pm range: two experiments). A dash indicates no statistically significant reduction in migration (all combinations of chemoattractant and tripeptide have been tested. The tripeptide WVQ inhibited migration in response to all chemoattractants tested and for this tripeptide the numbers shown are the ED₅₀ for the inhibition (mean of at least two determinations). Note that none of the tripeptides shown inhibited TGF- β 1 induced migration at 100 μ M. The bolded values indicate the inhibition by each peptide of migration, induced by the chemoattractant from which it was derived, i.e., KQK was derived from MCP-1, etc.

a The affinity constant for KQK binding to DARC is 15 μ M.
b The ED₅₀ for KQK inhibiting MCP-1 induced migration is 7 μ M.
c The affinity constant for WVQ binding to DARC is 2 μ M.

25

Example 8

In Vivo Pharmacokinetics and Toxicity

When ³H-D-ala peptide 3(1-12)[MCP-1] (³H-D-ala was attached to Asp) was given as an IV or SQ bolus to mice, peak serum concentrations were reached within 1 hour. This radiolabeled peptide was rapidly excreted (approximately 4 hours), primarily via the kidney. Biodistribution data indicated that the primary target organ was the kidney with much smaller amounts detected in blood, liver and intestine. In contrast, CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] was detected for 24 hours or more in the circulation, presumably as a result of its Duffy binding. Direct

comparison of ^3H -D-ala peptide 3(1-12)[MCP-1] (no DARC binding and rapidly cleared) and CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] (weak Duffy binding and good serum half-life) indicates that peptides of the invention may be particularly useful to increase the half-life of other pharmaceutical agents.

5 A modified LD₅₀ technique was used to determine the mouse intravenous LD₅₀ value for CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1]. The LD₅₀ = 11.4 mg/mouse IV, which is 569 mg/kg. This is equivalent to a human IV dose of 39 grams. This is ten times more than the efficacious dose seen in either the asthma model or the endotoxemia model (see Examples below). Intraperitoneal
10 administration of 11 mg did not result in lethality. Histologically, toxicity was confined to the kidneys and lymphoid tissues.

At the lethal dose, apoptosis of lymphocytes was seen in the spleen and gut-associated lymphoid tissue. The rate limiting toxicity was to the kidney. There was a dose dependent increase in acute tubular nephrosis. This is most likely due to the
15 huge intravenous bolus (569 mg/kg) of a small molecular weight peptide which is excreted very rapidly (first pass) by the kidney. This is very similar to the change seen in patients with massive release of myoglobin or hemoglobin after crush injuries or massive hemolysis. At the lethal dose, histologic evidence of acute tubular nephrosis and mild lymphoid cell death were seen.

20 Using an in-life phase of an acute rat toxicity study, no clinically detectable changes associated with test agent administration of doses up to 10 mg IV were found. In a 7-day repeat dose toxicity study in rats, no clinical signs were observed in treated animals.

25

Example 9

Use of the Agents of the Invention in a Rat Dermal Inflammation Model

To assess the efficacy of an agent of the invention in the prevention of lipopolysaccharide (LPS)- and MCP-1-induced dermal inflammation in the rat, three different doses of CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] were administered.

An inflammatory response was elicited by intradermal injection (ventral abdomen) of either 500 ng MCP-1 or 100 ng MCP-1 along with endotoxin-free phosphate-buffered saline vehicle (as a negative control) and bacterial lipopolysaccharide (LPS; as a positive control). Each substance was injected at a different site. The 5 results obtained from animals was compared to CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] treated, and PBS (diluent control) treated, animals. Thirty minutes prior to intradermal agonist administration, the animals received an intravenous loading dose (3, 30 or 300 mg) and a subcutaneous depo dose (0.1, 1 or 10 mg) (on dorsum) of the pan-chemokine inhibitor CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-10] (see, for example, Figure 17). Animals were sacrificed at 20-24 hours post injection. Serum and urine were collected. The intradermal sites of agonist injection were collected, bisected and the extent of the inflammatory response was assessed by histopathology and quantitative immunofluorescence (fixed and frozen) (for example, following MCP-1 injection, the number of monocyte/macrophages in 15 the skin was determined using the anti-CD14 (MCA342 from Serotec; clone ED2) at 3 µg/ml overnight at 4°C. The second antibody was rat anti-mouse FITC (415-096-100 from Jackson ImmunoResearch) at 28 µg/ml for 6 hours at room temperature). In addition, toxicity of CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] was assessed by collection of the following tissue samples in 10% neutral buffered formalin for 20 histologic analysis: lung, liver, kidney, spleen, thymus, heart, and antagonist (test agent) injection site.

The results of a typical experiment are shown in Figures 9 and 10. Systemic treatment with CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] completely abolished MCP-1 induced recruitment of monocyte/macrophages ($p = 0.009$). This is 25 consistent with potent inhibition of MCP-1-induced migration seen *in vitro* with this agent. Furthermore, CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] reduced the number of resident tissue monocyte/macrophages in the site that received PBS alone, and also in untreated skin. This is consistent with a systemic downregulation of monocyte/macrophage recruitment 24 hours after a single treatment with CRD-

Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1]. In contrast, D-ala peptide 3(1-12)[MCP-1] had no effect *in vivo* ($p = 0.754$), in accord with its lack of *in vitro* activity in the migration assay.

A substantial reduction (>80%) in the number of monocyte/macrophages recruited in response to injected bacterial LPS was also noted. LPS was a stronger inducer of macrophage recruitment than MCP-1 even at 500 ng dose. Previous studies suggested that LPS-mediated macrophage accumulation was heavily dependent on TNF- α (a non-chemokine chemoattractant) since neutralizing antibodies to TNF- α markedly reduced LPS-induced inflammation. However, in endotoxemia models (Example 10) CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] markedly reduced LPS-induced increases in plasma TNF- α suggesting that chemokines may play a role in the induction of TNF- α , and that both chemokine signaling and TNF- α signaling may be necessary for maximal LPS-induced inflammation.

Although MCP-1 is fairly specific as a monocyte/macrophage chemoattractant, dermal injection of LPS induces recruitment of a broader range of leukocytes, including T- and B-cells and neutrophils. Specific antibodies to rat B-cells (MCA 1432 from Serotec) were used at 10 μ g/ml overnight at 4°C to determine whether CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] affected the recruitment of this leukocyte subpopulation. Secondary antibody was anti-mouse FITC (415-096-100 from Jackson ImmunoResearch, as above). As for monocyte/macrophages, CRD-Leu₄Ile₁₁Cys₃ peptide 3(3-12)[MCP-1] substantially inhibited the recruitment of B-cells to the site of the LPS injection (Figure 10). Thus, the anti-inflammatory effects of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] and other peptide 3 derivatives, analogs and variants are not limited to reducing or inhibiting macrophage accumulation but also inhibit recruitment of other leukocyte subsets.

Example 10

Use of the Agents of the Invention in a Murine Endotoxemia Model

A mouse endotoxemia model is used to screen peptides for *in vivo* functional and cytokine activity in a rapid manner. Female CD-1 mice are injected i.p. (Ventral abdomen) with 583 µg LPS and TNF- α , IFN- γ , IL-4 and MCP-1 protein and mRNA levels determined. Thirty minutes prior to LPS administration, the animals were 5 administered one of three different doses of CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] as an intravenous loading dose and a subcutaneous bolus dose (on dorsum). PBS treated animals with and without LPS administration were positive 10 and negative controls. Two hours later, animals were euthanized and serum collected. Serum was separated from the cell pellet and frozen until ELISA analysis 15 of cytokine levels. Lung and liver samples were collected for mRNA analyses and histopathology. CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1] demonstrated a dose-dependent decrease in serum TNF- α . Serum levels and mRNA levels of IL-4, IFN- γ and MCP-1 are also determined.

Example 11

Use of the Agents of the Invention in a Mouse Asthma Model

To determine whether increasing doses of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] alters the cell number and type of cell within the lung, mice were 20 injected intravenously, intravenously and intratracheally, or intratracheally alone with CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1]. Mice were sacrificed at 20-24 hours post injection. Lungs were collected for isolation of cells, which were subsequently counted and characterized by surface staining for CD3, CD4, CD8, 25 B220, and Mac-1.

The total number of cells isolated from the lungs was higher in all groups receiving low dose CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] (0.3 µg IV and or 10 µg IT) compared to PBS-treated mice. There were no significant differences in

the total number of cells isolated from lungs of mice treated with the high dose CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] compared to PBS controls.

By FACS analysis, high dose CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] significantly reduced the percentages of CD3, CD4, and B220 cells by all routes of 5 administration compared to PBS controls. In contrast, there were not significant differences in the percentages of CD3, CD4, or B220 cells in the groups treated with low dose CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] by all routes of administration.

Another study assessed the ability of two increasing doses of CRD-
10 Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] to reduce the pulmonary inflammatory infiltrate, inhibit IgE antibody increases, and alter the percentages of specific inflammatory cells in the lung and blood in mice challenged intratracheally with ovalbumin. See Gonzalo et al., J. Clin. Invest., 98, 2332 (1996); Gonzalo et al., J. Exp. Med., 188, 157 (1998).

15 Mice were sensitized with 0.1 mg of ovalbumin in 200 μ l PBS (diluent control) intraperitoneally (Table 76). Eight days following sensitization, mice received an intravenous loading dose (0.3 or 30 μ g) and a subcutaneous depo dose (10 μ g or 1 mg) of the pan-chemokine inhibitor CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1]. Thirty minutes following CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-
20 1] administration, mice were challenged with 1% ovalbumin or PBS (diluent control) intratracheally. Twenty-one days following sensitization, mice received a second intravenous loading dose (0.3 or 30 μ g) and a subcutaneous dose (10 μ g or 1 mg) of the pan-chemokine inhibitor CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1]. Thirty minutes following CRD-Leu₄Ile₁₁Cys₁₃peptide 3(3-12)[MCP-1]
25 administration, mice were challenged with 2% ovalbumin or PBS (diluent control) intratracheally. Mice were sacrificed 3 hours post-ovalbumin challenge on day 21. Lungs were collected for histopathology and for isolation of cells for total cell counts and FACS analysis. PBLs were collected for FACS analysis. Serum was collected for IgE levels.

By FACS analysis, there were significantly lower percentages of CD3, CD4, B220, and Mac-1 cells in the lungs of mice treated with both doses of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] (0.3 IV/10 µg subcutaneously or 30 µg IV/1 mg subcutaneously) compared to mice which received PBS prior to challenge 5 with OVA. The percentage of CD8 cells was similar in all groups. In addition, the total number of cells isolated from lungs of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] mice was similar to PBS-treated mice but significantly lower than mice treated with OVA and PBS, suggesting that the agent altered trafficking of inflammatory cells into the lung. In the blood, there were significantly higher 10 percentages of CD3 and CD4 cells and lower percentages of B220 in mice treated with both doses of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] compared to OVA-treated mice (positive control) and to PBS-treated mice (diluent control). Mice treated with the high dose had fewer Mac-1 cells in the PBL compartment compared to all other groups.

15 Histologically, all mice treated with the high dose CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] had minimal to no inflammatory infiltrates in the lung, similar to mice treated with PBS alone. Mice that received low dose CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] also had minimal inflammation compared to mice treated with PBS and OVA. Rare eosinophils were seen only in the PBS OVA 20 group (positive control) which is an expected response to OVA sensitization.

IgE levels were significantly higher in mice treated with PBS and OVA compared to all other groups. IgE was not detectable above background in all groups of mice treated with CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1].

A third study assessed the ability of three increasing doses of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] to reduce the pulmonary inflammatory infiltrate, inhibit IgE antibody increases, and to alter the percentages of specific inflammatory cells in the lung of mice challenged intratracheally with ovalbumin. 25 Mice were sensitized with 0.1 mg of ovalbumin or PBS (diluent control) intraperitoneally. Eight days following sensitization, mice received a subcutaneous

dose (10.3 μ g, 103 μ g, or 1.03 mg) of the pan-chemokine inhibitor CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1]. Thirty minutes following CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] administration, mice were challenged with 1% ovalbumin or PBS (diluent control) intratracheally. Fifteen, eighteen, and 5 twenty-one days following sensitization, mice received subcutaneous doses (10.3 μ g, or 103 μ g, or 1.03 mg) of the pan-chemokine inhibitor CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1]. Thirty minutes following CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] administration, mice were challenged with 2% ovalbumin challenge, on day 21. Lungs were collected for histopathology and for isolation of 10 cells for total cell counts and FACS analysis. Serum was collected for IgE, IL-4, and IFN- γ levels.

By FACS analysis, there were significantly lower percentages of Mac-1 cells in the lung of mice treated with all doses of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] compared to mice which received PBS only prior to challenged with 15 OVA. Histologically, all mice treated with the high or medium dose CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] had fewer inflammatory infiltrates in the lung compared to mice that were not treated with the peptide but challenged with OVA (positive control). Mice treated with PBS alone had minimal to no 20 inflammation in the lung. All mice challenged with OVA had eosinophils in the lung. Similar to mice treated with PBS only (negative control), IgE levels were significantly lower in mice treated with CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] compared to mice treated with PBS and OVA (positive control).

Thus, CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1], when delivered IV and subcutaneously, or subcutaneously alone, altered the trafficking of lymphocytes into 25 the lung following exposure to an antigen. More significantly, CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] reduced the cellular inflammation in the lung, IgE responses and IL-4 concentration in the serum, which are strongly associated with asthma. IgE responses are dependent on a Th2 T cell response, which produces IL-4 and IL-5. Therefore, the observation that CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-

12)[MCP-1] has an effect on reducing IgE upon challenge with OVA strongly indicates that CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1] may reduce IL-4 and IL-5 also.

5

Example 12

Preferred Tripeptides and Analogs Thereof

Preferred tripeptides of the invention include KXK peptides, where X is one of the twenty naturally occurring amino acids, e.g., KQK and KLK, as well as peptides having KXK. As described below, KXK peptides are anti-inflammatory 10 by two distinct mechanisms. Some KXK peptides are TGF-beta activators and others are chemokine antagonists, and a subset are both (see Table 8).

Table 8

Peptide	TGF-beta Activator	Chemokine antagonist
KAK	n.d.	+
KCK	n.d.	n.d.
5 KDK	-	+
KEK	-	+
KFK	++++	-
KGK	-	-
10 KHK	n.d.	n.d.
KIK	++	++
KKK	-	++++
KLK	+++	++++
KMK	n.d.	n.d.
15 KNK	n.d.	+++
KPK	n.d.	n.d.
KQK	-	+++++
KRK	n.d.	n.d.
18 KSK	-	-
KTK	-	-
20 KVK	n.d.	n.d.
KWK	-	-
KYK	++++	-

To test whether a KXK tripeptide activates TGF-beta, a direct ELISA-type
25 assay can be used. Recombinant human latent TGF- β 1 produced in CHO cells
(R&D Systems) was incubated with the test activator. For example, 200 ng of latent
TGF- β 1 (at 20 μ g/ml) was incubated with test peptide at 100 nM final concentration
at 37°C for 90 minutes. Following incubation, the TGF- β is incubated with the
recombinant extracellular domain of the Type II TGF- β receptor (R2X) which binds
30 only the active and not the latent forms of TGF- β 1 (Clin. Chim. Acta, 235, 11
(1995)). For example, 1 μ g of purified R2X is coated onto a Maxisorp ELISA plate

well in 50 μ l of 100 mM sodium carbonate for 2 hours at 4°C, and non-specific protein binding then blocked incubation with 5% sucrose 5% Tween-20 in Tris-buffered saline for 1 hour at room temperature.

The TGF- β sample is then incubated with the coated and blocked wells for 2 hours at room temperature with shaking. Wells are washed 3 times quickly with Tris-buffered saline containing 0.05% Tween-20 between each incubation. If any of the latent TGF- β 1 has been activated by the incubation with test peptide, it is captured by the R2X, while remaining latent TGF- β 1 is washed away. Captured active TGF- β 1 is then detected by incubation with a suitable detection agent, such as a peroxidase conjugated polyclonal anti-TGF-beta antibody. For example, the wells are incubated with 200 μ l of BDA19 chicken anti-TGF- β 1 antibody coupled to horseradish peroxidase for 90 minutes at room temperature with shaking. Any bound peroxidase is then detected using a suitable chromogenic substrate (e.g., K-BLUE TMB substrate solution). The amount of active TGF- β generated is estimated by interpolation of a standard curve constructed using known amounts of active TGF- β 1 (R&D Systems).

Chemokine antagonist activity may be determined using the THP-1 transwell migration assay described above in which the peptide is incubated in the top compartment with the cells while a chemokine is used as a chemoattractant in the lower compartment. Four chemokines were tested: IL-8; SDF-1 α ; MCP-1 and MIP1 α : pluses in Table 8 indicate that the peptide was active as an inhibitor of migration induced by at least one of these four chemoattractant chemokines. The number of pluses is a qualitative indicating of the activity of each peptide in each assay. A minus indicates no detectable activity in the assay, and n.d. indicates that no attempt to estimate the activity of the given peptide in this assay has been made to date.

KFK was as active as RFK. However, in marked contrast to previous reports, other members of the KXK series were also active as TGF- β activators. For

example, the KYK was more active than KFK. Thus, the substitution of arginine for lysine increases the range of amino acids at position 2 which activate TGF- β .

KLK and KIK are of particular interest, since these agents are dual-action anti-inflammatory molecules. These tripeptides are specific antagonists of the 5 SDF-1 α receptor CXCR4, and also activate TGF- β . Thus, KLK, KIK and their analogs and derivatives are therefore likely to be particular useful pharmaceutical agents for the prevention or treatment of a wide range of anti-inflammatory disorders.

For graft eosinophilia, such as that associated with acute transplant rejection, 10 a pan-chemokine inhibitor, or a selective inhibitor of eosinophil recruitment (such as KKK or an analog thereof), may be particularly beneficial. Such agents may be used alone or in conjunction with lower than normal doses of steroids, such as prednisolone, which are used currently to control acute rejection episodes. Severe side-effects are associated with the use of the highest dose of prednisolone (or other 15 steroids) used during acute rejection, and use of agents which reduce or abolish the need to give steroids would be particularly useful.

Analogs of the KXK peptides, e.g., analogs of KQK, are also envisioned. The central chain (in a compound of formula V with R7 as a substituent) is replaced by a general substituent R, where R is the side chain from any of the amino acids. 20 These analogs (for example, the general class of fluoroalkenes of a compound of formula (VI),) are useful for the treatment of a wide variety of diseases where activation of TGF- β and/or inhibition of chemokine signaling are desired. By selecting an appropriate member of this class of molecules, it is possible to engineer the desired properties of the molecule. Thus, selection of KYK analogs provides 25 powerful activation of TGF- β in the absence of chemokine inhibition, while analogs of KLK have both properties. Analogs of KQK have inhibitory action on one or more chemokine receptors but do not activate TGF- β .

Thus, KYK, its analogs, and derivatives may be selected for use in diseases where TGF- β upregulation is particularly beneficial, for example, in atherosclerosis

or osteoporosis. In contrast, analogs of KQK may be selected where chemokine inhibition is desired but TGF- β upregulation may not be beneficial, for example, in treatment of HIV infection.

The KXK peptides, and isosteres thereof, may be useful to treat low bone
5 mineral density, where TGF-beta elevation and selective inhibition of MCP-1 are likely to be especially synergistic.

Derivatives or analogs of the KXK class may be used alone or in combination with other therapies for the treatment of inflammatory disorders, or other diseases or indications such as those described herein. For example,
10 derivatives, or analogs of KYK may be used in conjunction with steroids for the treatment of inflammatory conditions, allowing lower than normal doses of steroids to be used reducing the side effects associated with chronic steroid use.

It is also envisioned that conservative substitutions of the amino acids at positions 1 and 3 do not affect the activities of the molecules. Thus, one or both of
15 the lysine side chains (either in a peptide or in an analog such as (VI)) may be substituted with an arginyl side chain or an ornithinyl side chain.

All publications, patents and patent applications are incorporated herein by reference. While in the foregoing specification this invention has been described in
20 relation to certain preferred embodiments thereof, and many details have been set forth for purposes of illustration, it will be apparent to those skilled in the art that the invention is susceptible to additional embodiments and that certain of the details described herein may be varied considerably without departing from the basic principles of the invention.

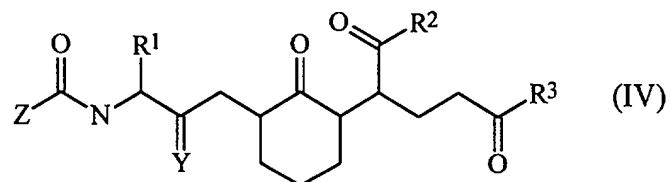
WHAT IS CLAIMED IS:

1. Chemokine peptide 3, a variant, or a derivative thereof.
2. Chemokine peptide 2, a variant, or a derivative thereof.
3. The peptide of claim 1 wherein the chemokine is not IL8 or NAP-2.
4. The peptide of claim 1 which is a variant of peptide 3[MCP-1].
5. The peptide of claim 4 which is Leu₄Ile₁₁peptide 3(3-12)[MCP-1].
6. The peptide of claim 1 or 2 which is a CC chemokine.
7. The peptide of claim 6 wherein the CC chemokine is MCP-1, RANTES, MCP-2, MCP-3, MCP-4, eotaxin, MIP1 α , MIP1 β , LARC, I309, HCC-1, TARC or Ck β 8.
8. The peptide of claim 1 or 2 which is a CXC chemokine.
9. The peptide of claim 8 wherein the CXC chemokine is IP-10, PF-4, SDF-1, NAP-2, GRO α , GRO β , GRO γ or ENA78.
10. The peptide of claim 8 wherein the CXC chemokine is IL-8, IP-10, SDF-1, PF-4, NAP-2, GRO α , GRO β , GRO γ , NAP-2 or ENA78.
11. A CRD derivative of chemokine peptide 3 or a variant thereof.

12. The derivative of claim 11 which is CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1].

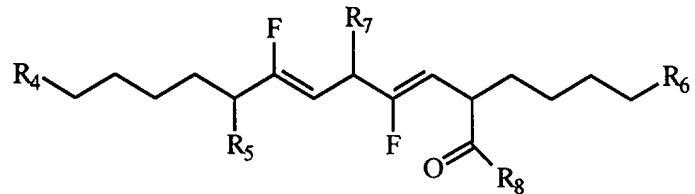
13. A CRD derivative of chemokine peptide 2 or a variant thereof.

14. A compound of formula (IV):



wherein R¹ is aryl, heteroaryl, coumaryl or chromanyl; wherein R² is N(R^a)(R^b); wherein R³ is N(R^c)(R^d); wherein Y is oxo or thioxo; wherein Z is (C₁-C₁₀)alkyl; wherein R^a-R^d are each independently hydrogen, (C₁-C₁₀)alkyl, (C₁-C₁₀)alkanoyl, phenyl, benzyl or phenethyl; or wherein R^a and R^b, or R^c and R^d, together with the nitrogen to which they are attached form a pyrrolidino, piperidino or morpholino ring; or a pharmaceutically acceptable salt thereof.

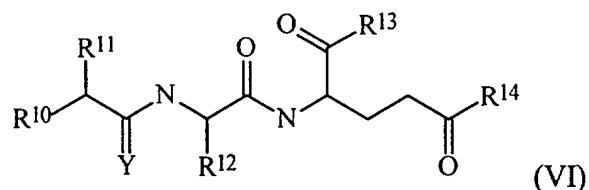
15. A compound of formula (V):



wherein R⁴ is NR_kR_l; wherein R⁵ is NR_mR_n; wherein R⁶ is NR_oR_p; wherein R⁷ is NR_qR_r; wherein R⁸ is hydrogen, hydroxy, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl,

(C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy, NR_sR_t, the N-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues; wherein R_k, R_l, R_o, and R_p are each hydrogen; wherein R_m and R_n are each independently hydrogen, acetyl, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, propoxy, butoxy, *tert*-butoxycarbonyl, 9-fluorenylmethoxycarbonyl, the C-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues; wherein R_q and R_r are each independently hydrogen, (C₁-C₁₀)alkyl, or (C₃-C₆)cycloalkyl; and wherein R_s are R_t are each independently hydrogen, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, phenyl, benzyl, or phenethyl; or a pharmaceutically acceptable salt thereof.

16. A compound of formula (VI):



wherein R¹⁰ is NRⁱR^j; R¹¹ is aryl, heteroaryl, aryl(C₁-C₃)alkyl, heteroaryl(C₁-C₃)alkyl, coumaryl, coumaryl(C₁-C₃)alkyl, chromanyl or chromanyl(C₁-C₃)alkyl; wherein any aryl or heteroaryl group, or the benz-ring of any coumaryl or chromanyl group may optionally be substituted with one, two or three substituents selected from the group consisting of halo, nitro, cyano, hydroxy, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, (C₁-C₆)alkanoyl, (C₂-C₆)alkanoyloxy, -C(=O)(C₁-C₆)alkoxy, C(=O)NR^eR^f, NR^eR^f; R¹² is (C₁-C₆)alkyl; R¹³ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy, hydroxy, or N(R^a)(R^b); R¹⁴ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or N(R^c)(R^d); Y is oxo or thioxo; and wherein R^a-R^j are each independently hydrogen, (C₁-C₁₀)alkyl, (C₁-C₁₀)alkanoyl,

phenyl, benzyl, or phenethyl; or R^a and R^b, R^c and R^d, R^e and R^f, R^g and R^h, or Rⁱ and R^j together with the nitrogen to which they are attached form a ring selected from pyrrolidino, piperidino, or morpholino; or a pharmaceutically acceptable salt thereof.

17. A method of preventing or inhibiting an indication associated with a chemokine-induced activity, comprising: administering to a mammal afflicted with, or at risk of, the indication an amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, or a combination thereof, effective to prevent or inhibit said activity, wherein the chemokine is not IL8 or NAP-2.
18. A method to inhibit the activity of more than one chemokine, comprising: administering to a mammal in need thereof an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.
19. A method to increase or enhance a chemokine-associated inflammatory response in a mammal, comprising: administering to the mammal an amount of a chemokine peptide 2, a variant thereof, a derivative thereof, or a combination thereof effective to increase or enhance said response.
20. A method of preventing or inhibiting an indication associated with hematopoietic cell recruitment, comprising: administering to a mammal at risk of, or afflicted with, the indication an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

21. A method of preventing or inhibiting an indication associated with histamine release from basophils or mast cells, comprising administering to a mammal at risk of, or afflicted with, the indication an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

22. A method to modulate the chemokine-induced activity of hematopoietic cells at a preselected physiological site, comprising: administering to a mammal a dosage form comprising an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof, wherein the dosage form is linked to a site targeting moiety.

23. A method to augment an immune response, comprising: administering to a mammal an immunogenic moiety and an amount of a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof, wherein the amount is effective to augment the immune response of the mammal to the immunogenic moiety.

24. A therapeutic method to prevent or treat a vascular indication, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof, wherein the indication is coronary artery disease, myocardial infarction, unstable angina pectoris, atherosclerosis or vasculitis.

25. A therapeutic method to prevent or inhibit lentiviral infection or replication, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

26. The method of claim 25 wherein the lentivirus is HIV.

27. The method of claim 26 further comprising administering an antiviral agent before, during and/or after the administration of the peptide, a variant thereof, derivative thereof, the compound of formula (IV), the compound of formula (V), the compound of formula (VI), or a combination thereof.

28. A therapeutic method to prevent or treat low bone mineral density, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

29. A method of inhibiting a parasitic infection in a vertebrate animal, comprising: administering to the animal an effective amount of a chemokine peptide 2, a variant thereof, a derivative thereof, or a combination thereof.

30. The method of claim 29 wherein the animal is a human with malaria.

31. A therapeutic method to prevent or treat an autoimmune disease, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative

thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

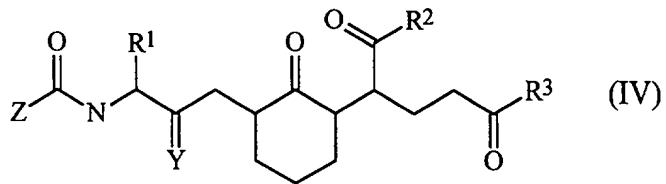
32. A method of suppressing tumor growth in a vertebrate animal, comprising: administering to said vertebrate an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

33. A method for preventing or treating psoriasis in a mammal, comprising: administering to the mammal an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

34. A method to increase or enhance hematopoietic cell-associated activity at a tumor site, comprising: administering an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

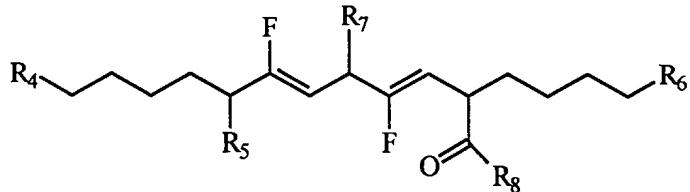
35. A method to enhance wound healing, comprising: administering an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

36. A method of treating a mammal afflicted with, or at risk of, an indication associated with chemokine-induced activity, comprising: administering to the mammal an effective amount of a compound of formula (IV):



wherein R^1 is aryl, heteroaryl, coumaryl or chromanyl; wherein R^2 is $N(R^a)(R^b)$; wherein R^3 is $N(R^c)(R^d)$; wherein Y is oxo or thioxo; wherein Z is (C_1-C_{10}) alkyl; wherein R^a-R^d are each independently hydrogen, (C_1-C_{10}) alkyl, (C_1-C_{10}) alkanoyl, phenyl, benzyl or phenethyl; or wherein R^a and R^b , or R^c and R^d , together with the nitrogen to which they are attached form a pyrrolidino, piperidino or morpholino ring; or a pharmaceutically acceptable salt thereof.

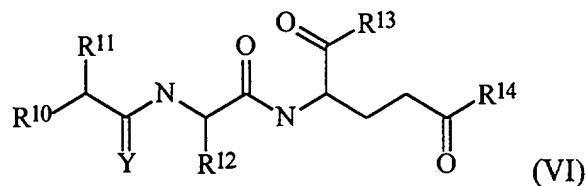
37. A method of treating a mammal afflicted with, or at risk of, an indication associated with chemokine-induced activity, comprising: administering to the mammal an effective amount of a compound of formula (V):



wherein R^4 is NR_kR_l ; wherein R^5 is NR_mR_n ; wherein R^6 is NR_oR_p ; wherein R^7 is NR_qR_r ; wherein R^8 is hydrogen, hydroxy, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, (C_3-C_6) cycloalkyl(C_1-C_6)alkyl, (C_1-C_{10}) alkoxy, (C_3-C_6) cycloalkyl(C_1-C_6)alkoxy, NR_sR_t , the N-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues; wherein R_k , R_l , R_o , and R_p are each hydrogen; wherein R_m are R_n are each independently hydrogen, acetyl, (C_1-C_{10}) alkyl, (C_3-C_6) cycloalkyl, propoxy, butoxy,

tert-butoxycarbonyl, 9-fluorenylmethoxycarbonyl or the C-terminal residue of an amino acid or a peptide of 2 to about 25 amino acid residues; wherein R_q and R_r are each independently hydrogen, (C₁-C₁₀)alkyl, or (C₃-C₆)cycloalkyl; and wherein R_s are R_t are each independently hydrogen, (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, phenyl, benzyl, or phenethyl; or a pharmaceutically acceptable salt thereof.

38. A method of treating a mammal afflicted with, or at risk of, an indication associated with chemokine-induced activity, comprising: administering to the mammal an effective amount of a compound of formula (VI):



wherein R¹⁰ is NRⁱR^j; R¹¹ is aryl, heteroaryl, aryl(C₁-C₃)alkyl, heteroaryl(C₁-C₃)alkyl, coumaryl, coumaryl(C₁-C₃)alkyl, chromanyl or chromanyl(C₁-C₃)alkyl; wherein any aryl or heteroaryl group, or the benz-ring of any coumaryl or chromanyl group may optionally be substituted with one, two or three substituents selected from the group consisting of halo, nitro, cyano, hydroxy, (C₁-C₆)alkyl, (C₁-C₆)alkoxy, (C₁-C₆)alkanoyl, (C₂-C₆)alkanoyloxy, -C(=O)(C₁-C₆)alkoxy, C(=O)NR^gR^h, NR^eR^f; R¹² is (C₁-C₆)alkyl; R¹³ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy, hydroxy, or N(R^a)(R^b); R¹⁴ is (C₁-C₁₀)alkyl, (C₃-C₆)cycloalkyl, (C₃-C₆)cycloalkyl(C₁-C₆)alkyl, (C₁-C₁₀)alkoxy, (C₃-C₆)cycloalkyl(C₁-C₆)alkoxy or N(R^c)(R^d); Y is oxo or thioxo; and wherein R^a-R^j are each independently hydrogen, (C₁-C₁₀)alkyl, (C₁-C₁₀)alkanoyl, phenyl, benzyl, or phenethyl; or R^a and R^b, R^c and R^d, R^e and R^f, R^g and R^h or Rⁱ and R^j together with the nitrogen to which they are attached form a ring

selected from pyrrolidino, piperidino, or morpholino; or a pharmaceutically acceptable salt thereof.

39. An immunogenic composition comprising an immunogenic moiety and an amount of a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

40. A therapeutic method to prevent or treat asthma, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

41. The method of claim 17 wherein the amount inhibits a product or intermediate in the arachidonic acid pathway.

42. The method of claim 41 wherein leukotriene is inhibited.

43. The method of claim 41 wherein thromboxane is inhibited.

44. The method of claim 41 wherein prostaglandin is inhibited.

45. A method of preventing or inhibiting an indication associated with elevated TNF- α , comprising: administering to a mammal afflicted with, or at risk of, the indication an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI) or a combination thereof.

46. A peptide that includes the amino acid sequence KXX, which is a chemokine antagonist, activates TGF-beta, or a combination thereof.

47. CRD-Cys₁₃Leu₄Ile₁₁peptide 3(3-12)[MCP-1].

48. A therapeutic method to prevent or treat organ transplant rejection, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

49. A therapeutic method to prevent or treat rheumatoid arthritis, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), a compound of formula (VI), or a combination thereof.

50. A therapeutic method to prevent or treat allergy, comprising: administering to a mammal in need of such therapy an effective amount of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), or a compound of formula (VI), or a combination thereof.

51. Use of a chemokine peptide 3, a chemokine peptide 2, a variant thereof, a derivative thereof, a compound of formula (IV), a compound of formula (V), or a compound of formula (VI), or a combination thereof for the manufacture of a medicament for the treatment of a pathological condition or symptom in a mammal which is associated with a chemokine-induced activity.

a mammal which is associated with a chemokine-induced activity.

Abstract of the Invention

Isolated and purified chemokine peptides, variants, and derivatives thereof, as well as chemokine peptide analogs, are provided.

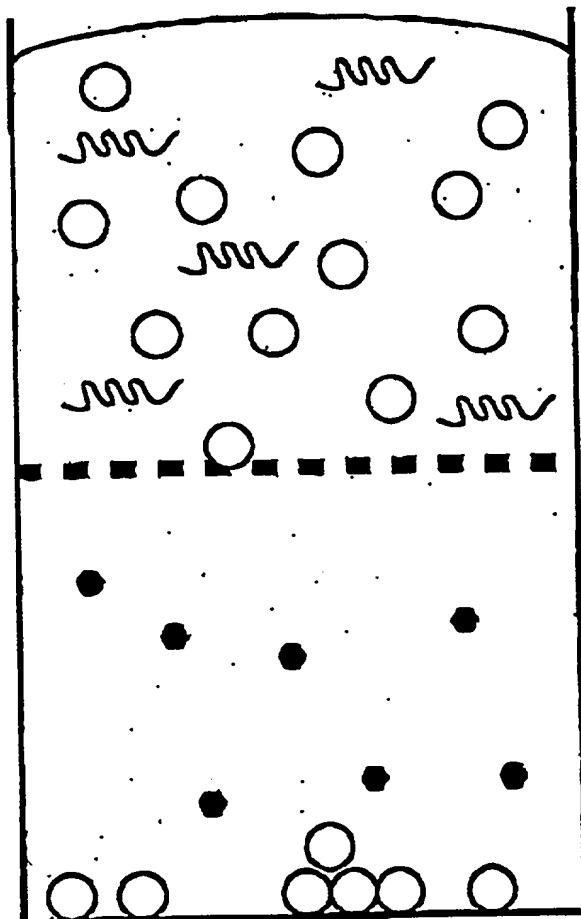
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Date of Deposit: September 11, 1998
I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 on the date indicated above and is addressed to the Assistant Commissioner for Patents, Washington, D.C. 20231

Printed Name JANET E. FIMMELSON

Signature Janet E. Fimmlson

TOP COMPARTMENT



LOWER COMPARTMENT

FIGURE 1

Effect of Peptide 3 with MCP-1 (50ng/ml)

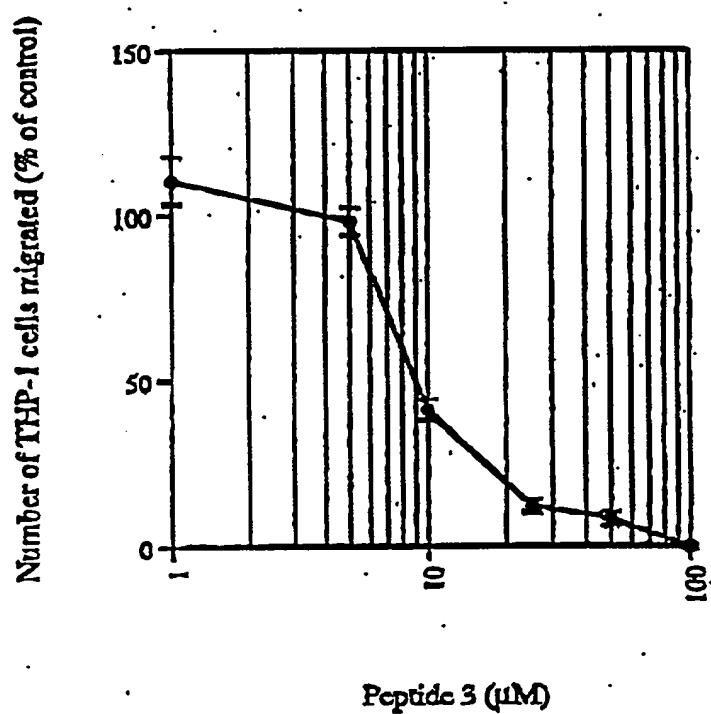


FIGURE 2

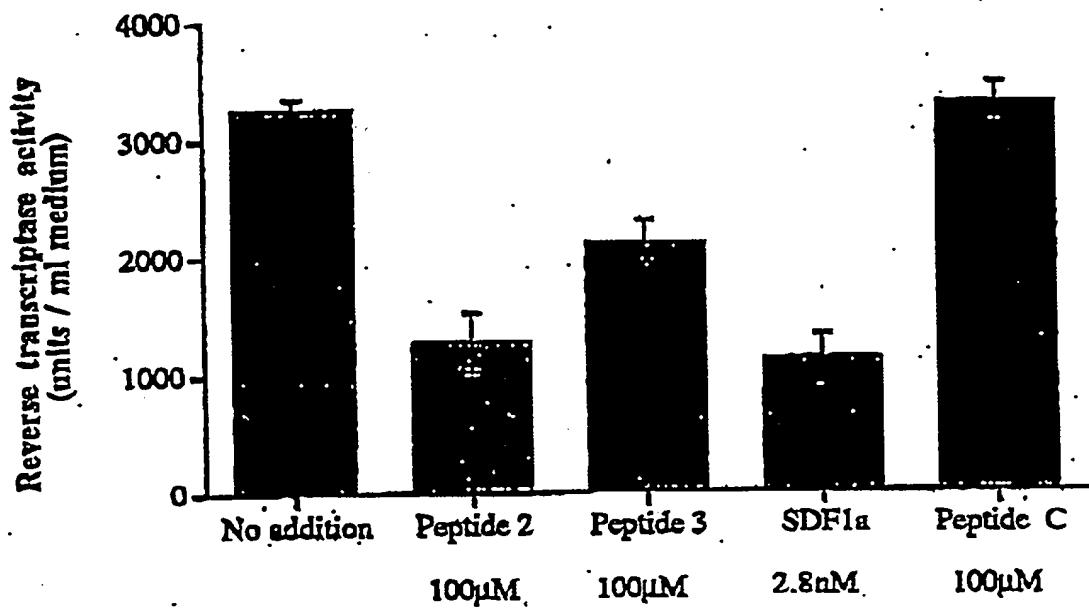


FIGURE 3

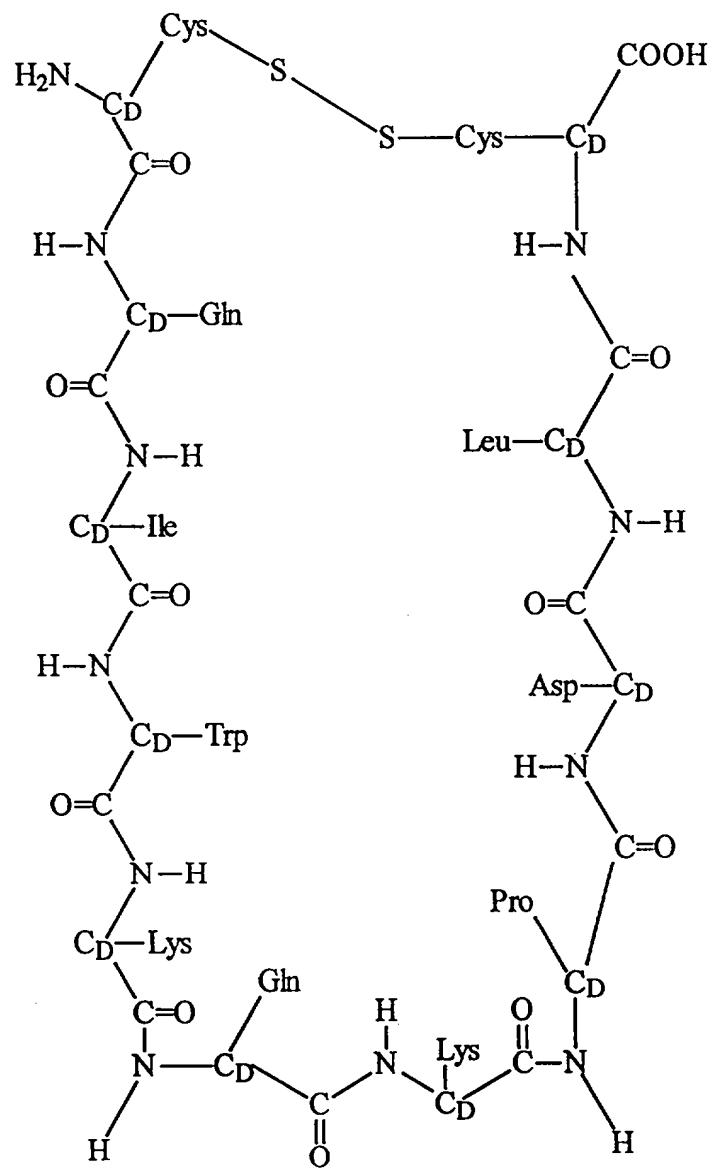
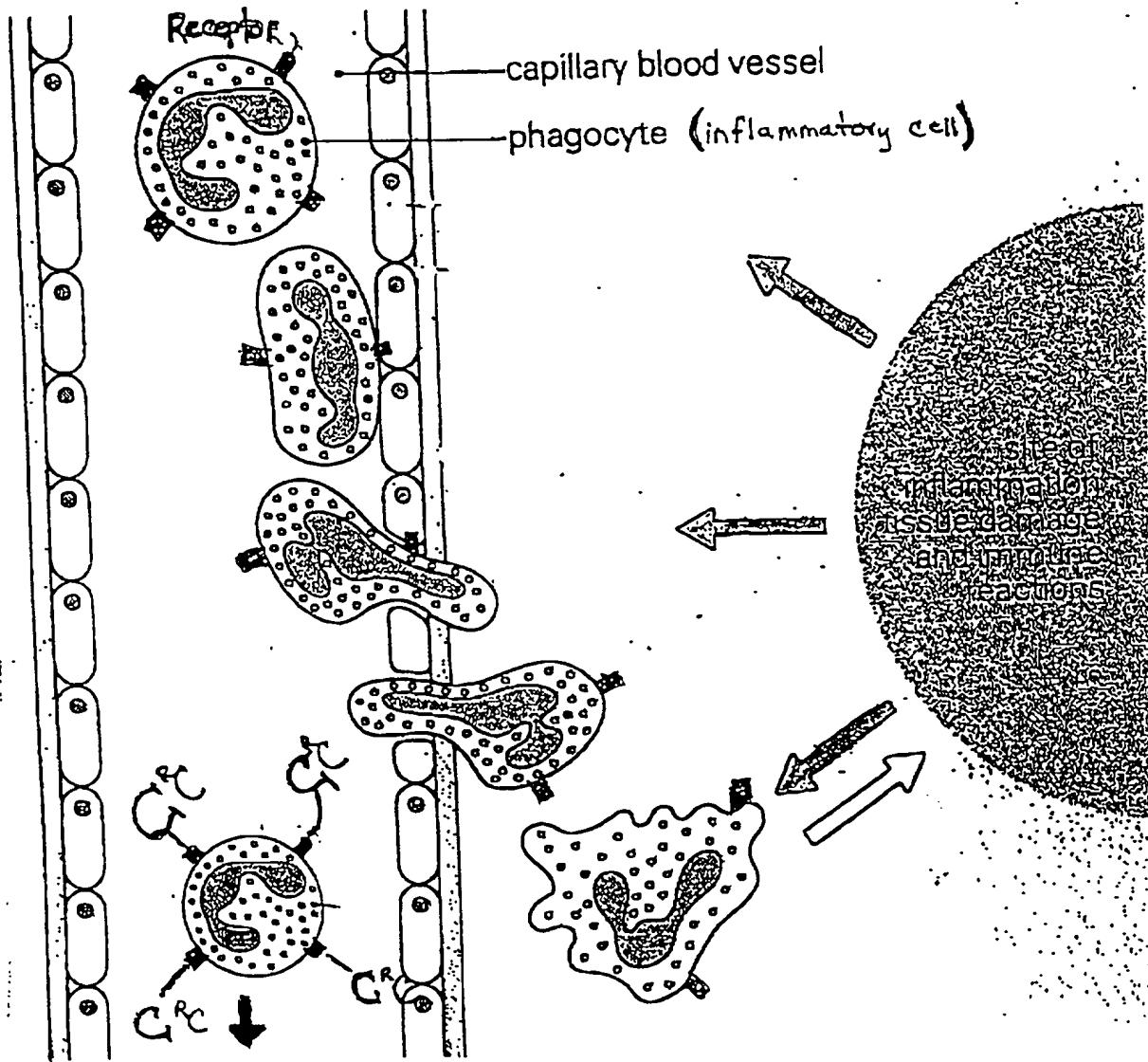


FIGURE 4



Figures

TARGET FOR VIRAL ENTRY (HIV VIRUS)

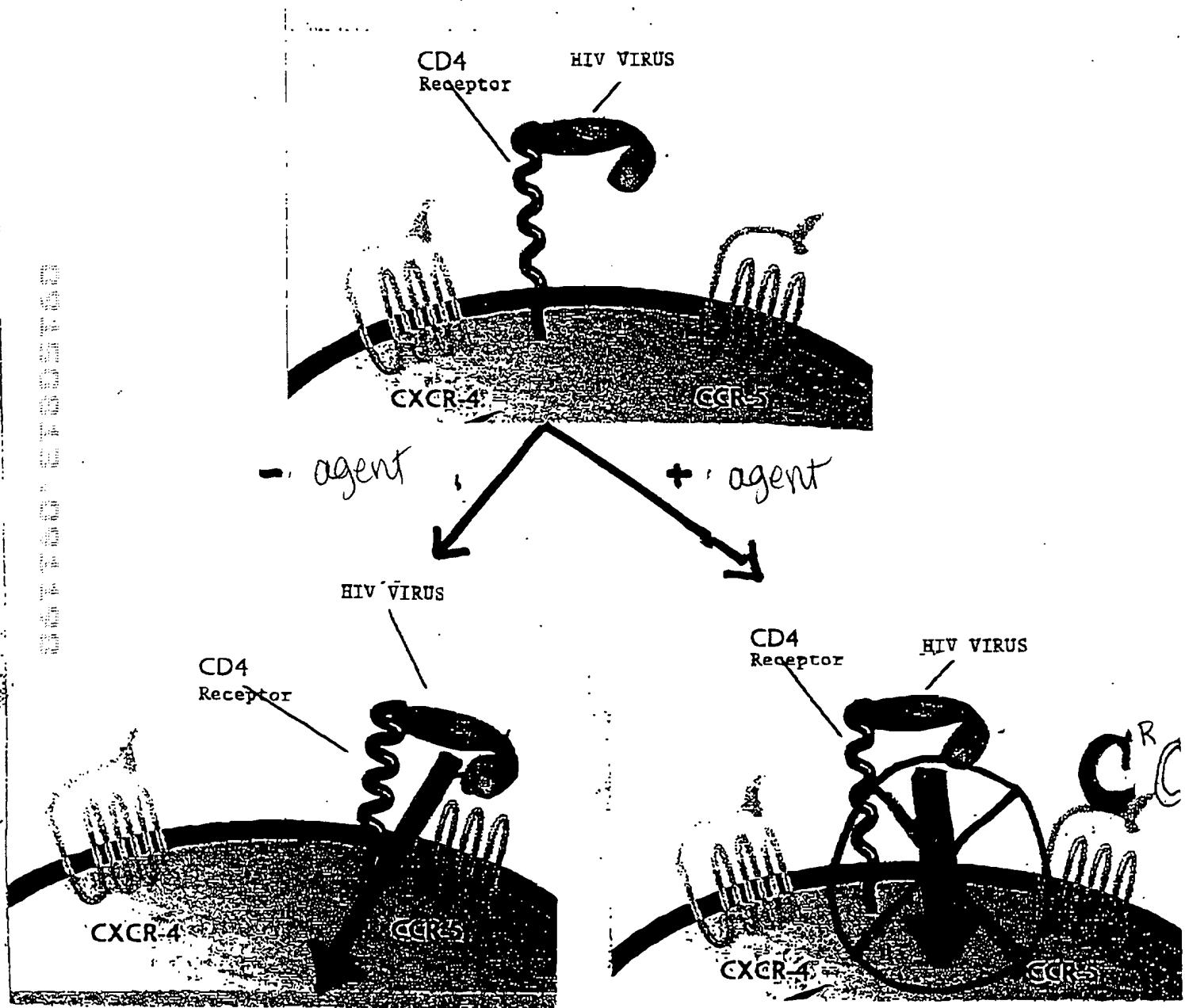
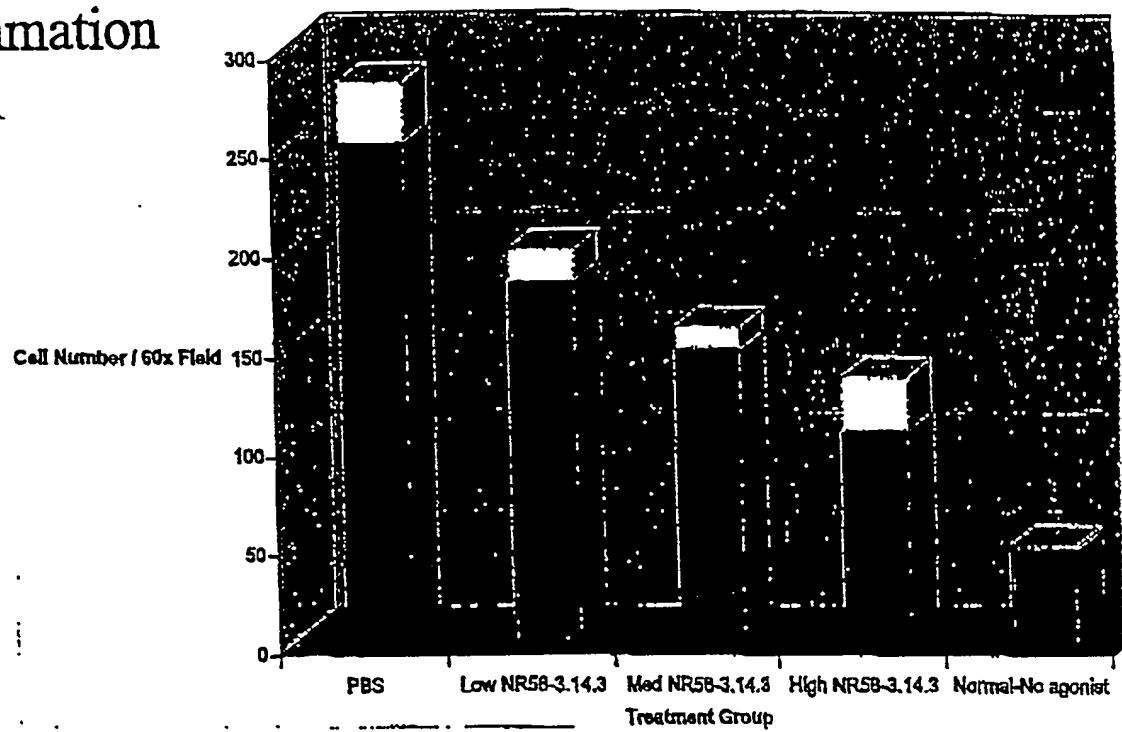


Figure 6

Skin Inflammation Model



Endotoxemia Model

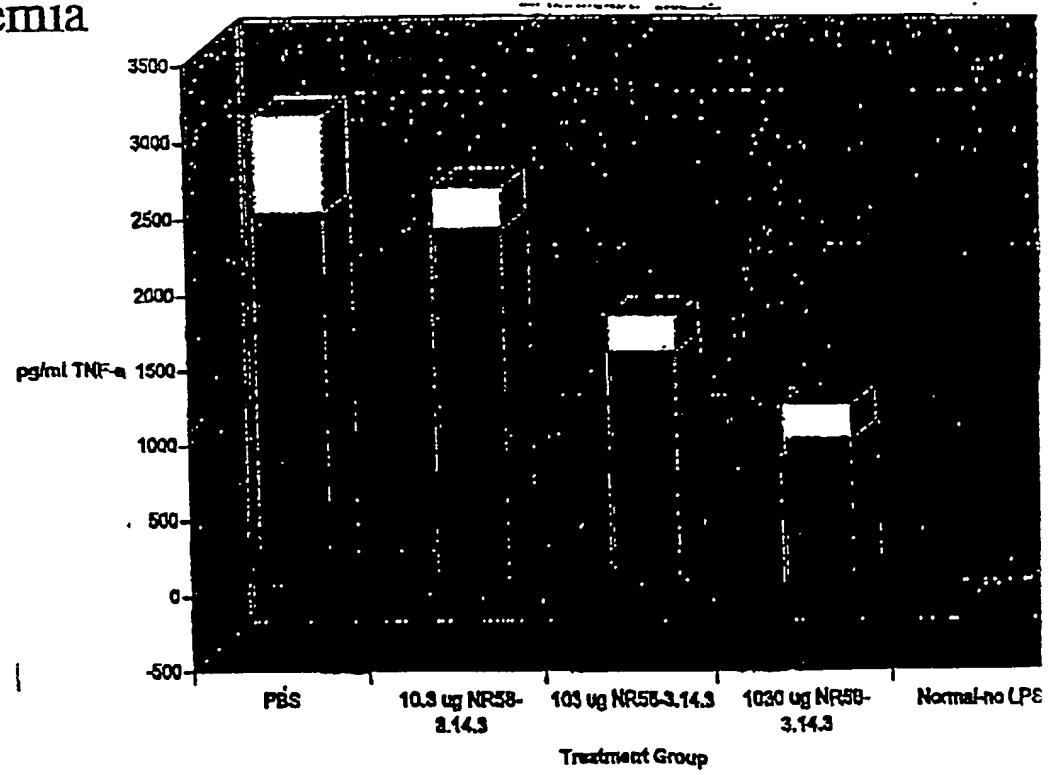


Figure 7

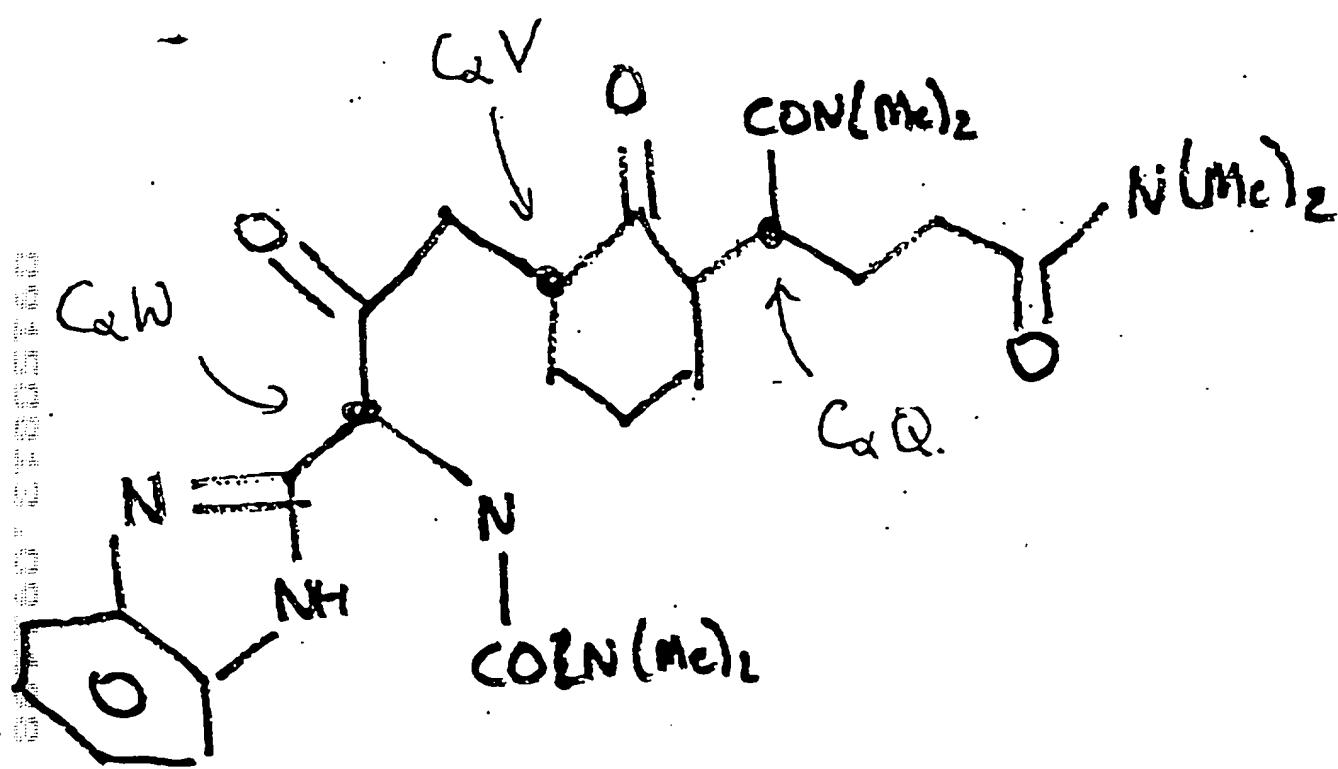


Figure 8

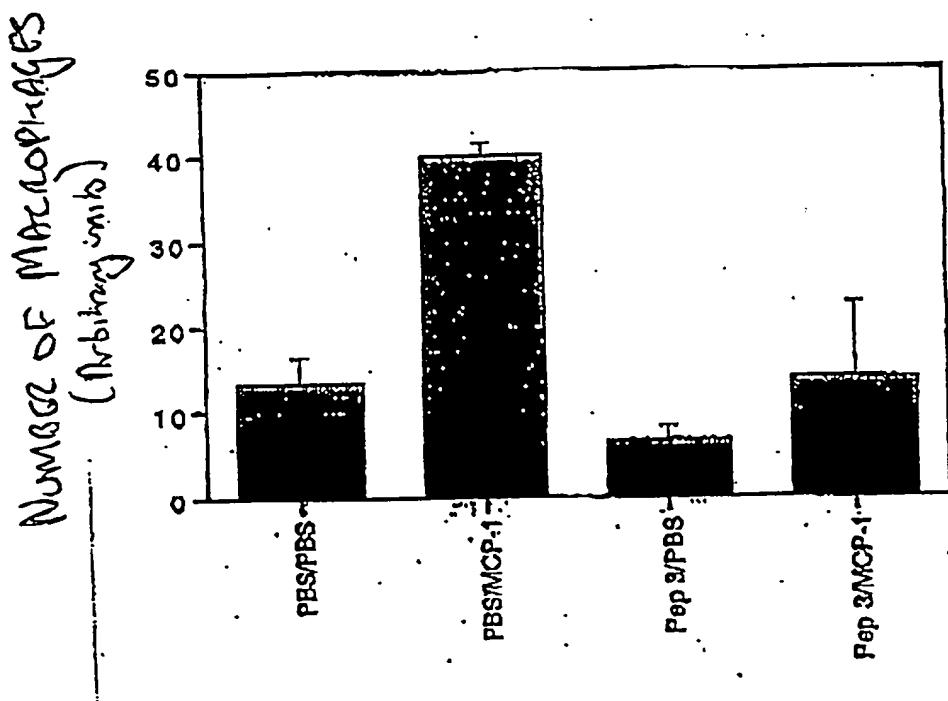


Figure X
9

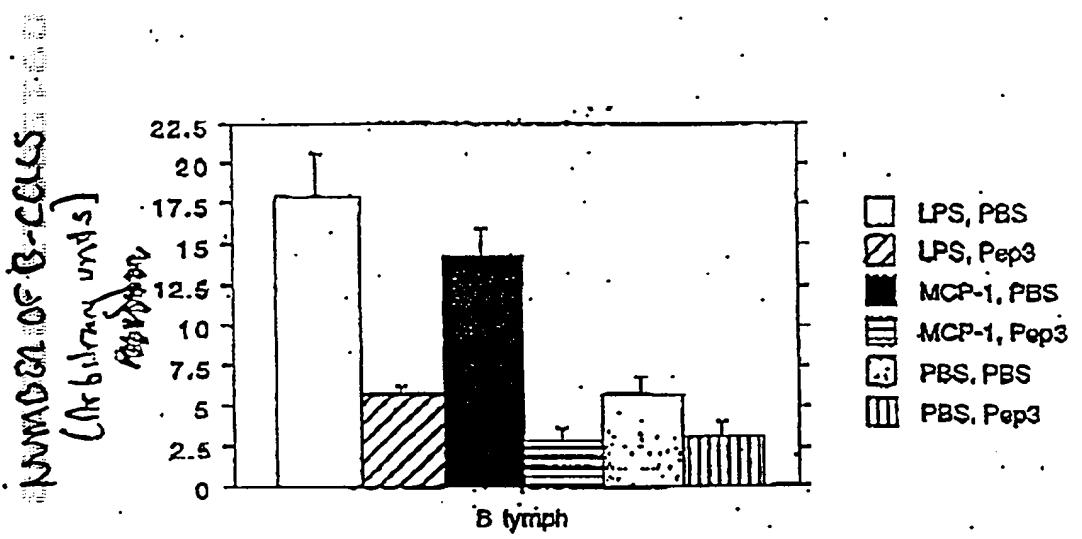


Figure X
10

THP-1 cells with M-Tropic HIV

FRACTION
OF
CELLS
INFECTED
(infected / total cells)

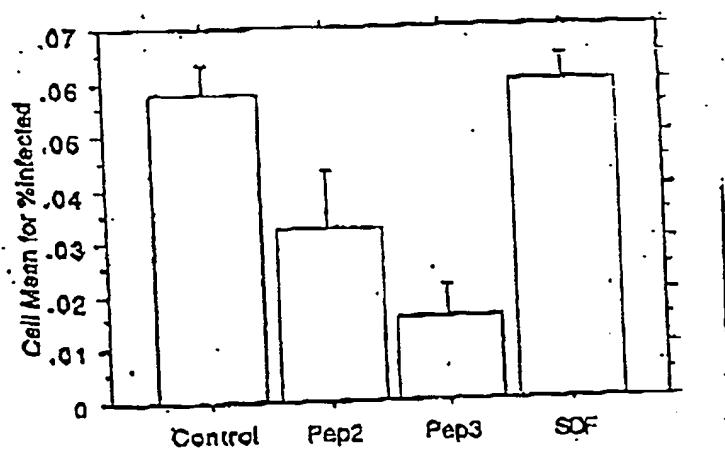


FIGURE B. 2 11

Figure 12

<u>Amino Acid</u>	<u>Codon</u>
Phe	UUU, UUC
Ser	UCU, UCC, UCA, UCG, AGU, AGC
Tyr	UAU, UAC
Cys	UGU, UGC
Leu	UUA, UUG, CUU, CUC, CUA, CUG
Trp	UGG
Pro	CCU, CCC, CCA, CCG
His	CAU, CAC
Arg	CGU, CGC, CGA, CGG, AGA, AGG
Gln	CAA, CAG
Ile	AUU, AUC, AUA
Thr	ACU, ACC, ACA, ACG
Asn	AAU, AAC
Lys	AAA, AAG
Met	AUG
Val	GUU, GUC, GUA, GUG
Ala	GCU, GCC, GCA, GCG
Asp	GAU, GAC
Gly	GGU, GGC, GGA, GGG
Glu	GAA, GAG

FIGURE 13

Original Residue	Exemplary Substitutions	Preferred Substitutions
Ala (A)	val; leu; ile	val
Arg (R)	lys; gln; asn	lys
Asn (N)	gln; his; lys; arg	gln
Asp (D)	glu	glu
Cys (C)	ser	ser
Gln (Q)	asn	asn
Glu (E)	asp	asp
Gly (G)	pro	pro
His (H)	asn; gln; lys; arg	arg
Ile (I)	leu; val; met; ala; phe norleucine	leu
Leu (L)	norleucine; ile; val; met; ala; phe	ile
Lys (K)	arg; gln; asn	arg
Met (M)	leu; phe; ile	leu
Phe (F)	leu; val; ile; ala	leu
Pro (P)	gly	gly
Ser (S)	thr	thr
Thr (T)	ser	ser
Trp (W)	tyr	tyr
Tyr (Y)	trp; phe; thr; ser	phe
Val (V)	ile; leu; met; phe; ala; norleucine	leu

FIGURE 14

Peptide 3

LFL peptide 3(1-12)[MCP-1]: Residues 50-61 of mature hMCP-1
E-I-C-A-D-P-K-Q-K-W-V-Q
L amino acids

LFL peptide 3(3-12)[MCPI] Residues 52-61 of mature hMCP-1
C-A-D-P-K-Q-K-W-V-Q
L amino acids

LFL peptide 3(1-6)[MCP1]: residues 50-55 of mature hMCP-1
E-I-C-A-D-P
L amino acids

LFL peptide 3(7-12)[MCP1]: Residues 56-61 of mature hMCP-1
K-Q-K-W-V-Q
L amino acids

LFL Leu₄peptide3(1-12)[MCP-1]
E-I-C-L-D-P-K-Q-K-W-V-Q
L amino acids

LFL Ser₇peptide3(1-12)[MCP-1]
E-I-C-A-D-P-S-Q-K-W-V-Q
L amino acids

LFL Ile₁₁peptide3(1-12)[MCP-1]
E-I-C-A-D-P-K-Q-K-W-I-Q
L amino acids

LFL Leu₄Ile₁₁peptide3(1-12)[MCP-1]
E-I-C-L-D-P-K-Q-K-W-I-Q
L amino acids

CFL Cys₀Leu₄Ile₁₁Cys₁₃peptide3(1-12)[MCP-1]
C-E-I-C-L-D-P-K-Q-K-W-I-Q-C
L amino acids

LRD Leu₄Ile₁₁ peptide 3(1-12)[MCP-1]
q-i-w-k-q-k-p-d-l-c-i-e
D amino acids

CRD Cys₀Leu₄Ile₁₁Cys₁₃peptide 3(1-12)[MCP-1]
c-q-i-w-k-q-k-p-d-l-c-i-e-c
D amino acids

LFL Ser₇Glu₈Glu₉peptide3(1-12)[MCP1]:Residues 50-61 of mature hMIP1 α
E-I-C-A-D-P-S-E-E-W-V-Q
L amino acids

LFL peptide3(10-12)[MCP-1]
W-V-Q
L amino acids

CFL Cys₀Cys₄ peptide3(10-12)[MCP-1]
C-W-V-Q-C
L amino acids

LRD peptide3(10-12)[MCP-1]
q-v-w
D amino acids

LFL peptide3(7-9)[MCP-1]
K-Q-K
L amino acids

LRD peptide3(7-9)[MCP-1]
k-q-k
D amino acids

LFL peptide 3(7-9)[MIP1 α](MIP1 α specific inhibitor)
S-E-E
L amino acids

LRD peptide3(7-9)[MIP1 α] (MIP1 α specific inhibitor)
e-e-s
D amino acids

LFL peptide3(7-9)[IL-8](IL-8 specific inhibitor)
K-E-N
L amino acids

LRD peptide3(7-9)[IL-8](IL-8 specific inhibitor)
n-e-k
D amino acids

LFL peptide3(7-9)[SDF-1 α](SDF-1 α specific inhibitor)
K-L-K
L amino acids

LRD peptide3(7-9)[SDF1 α] (SDF-1 α specific inhibitor)
k-l-k
D amino acids

LFL Leu₄Ile₁₁Cys₁₃ peptide3(3-12)[MCP-1]
L-D-P-K-Q-K-W-I-Q-C
L amino acids

CRD Leu₄Ile₁₁Cys₁₃ peptide3(3-12)[MCP-1]
c-q-i-w-k-q-k-p-d-l-c
D amino acids

³H-Ala CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1](D-Ala attached to Asp residue of CRD-Leu₄Ile₁₁Cys₁₃ peptide 3(3-12)[MCP-1])

³H-L-Leu LRD Cys₁₃ peptide3(3-12)[MCP-1]
c-q-i-w-k-q-k-p-d-L-c
D and L amino acids

LFL SES
S-E-S
L amino acids

LFL KKK
K-K-K
L amino acids

LFL Cys₄ peptide3(10-12)[MCP-1]
W-V-Q-C
L amino acids

LRD Cys₄ peptide3(10-12)[MCP-1]
c-q-v-w
D amino acids

LFL Ile₁₁Cys₁₃ peptide3(10-12)[MCP-1]
W-I-Q-C
L amino acids

LRD Cys₁₃Ile₁₁peptide3(10-12)[MCP-1]

cqiw

D amino acids

LRD peptide3(7-12)[MCP-1]

q-v-w-k-q-k

D amino acids

CFL Cys₀Cys₁₃peptide3(7-12)[MCP-1]

C-K-Q-K-W-V-Q-C

L amino acids

CRD Cys₀Cys₁₃peptide3(7-12)[MCP-1]

c-q-v-w-k-q-k-c

D amino acids

LFL peptide3(10-12)[RANTES]

WVR

L amino acids

LRD peptide3(10-12)[RANTES]

rvw

D amino acids

LFL peptide3(10-12)[SDF-1]

W-I-Q

L amino acids

Peptide 2

LFL peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1

S-Y-R-R-I-T-S-S-K-C-P-K-E-A-V

L amino acids

CFL Cys₀Cys₁₆peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1

C-S-Y-R-R-I-T-S-S-K-C-P-K-E-A-V-C

L amino acids

LRD peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1

v-a-e-k-p-c-k-s-s-t-i-r-r-y-s

D amino acids

CRD Cys₀Cys₁₆peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1
c-v-a-e-k-p-c-k-s-s-t-i-r-r-y-s-c
D amino acids

LFL peptide 2(1-15)[SDF1]: Residues 26-40 of mature hSDF-1 β
H-L-K-I-L-N-T-P-N-C-A-L-Q-I-V
L amino acids

CFL Cys₀Cys₁₆peptide 2(1-15)[SDF1]: Residues 26-40 of mature hSDF-1 β
C-H-L-K-I-L-N-T-P-N-C-A-L-Q-I-V-C
L amino acids

LRD peptide 2(1-15)[SDF1]: Residues 26-40 of mature hSDF-1 β
v-i-q-l-a-c-n-p-t-n-l-i-k-l-h
D amino acids

CRD Cys₀Cys₁₆peptide 2(1-15)[SDF1]: Residues 26-40 of mature hSDF-1 β
c-v-i-q-l-a-c-n-p-t-n-l-i-k-l-h-c
D amino acids

LFL peptide 2(1-14)[MIP-1 α]: Residues 28-41 of hMIP-1 α
D-Y-F-E-T-S-S-Q-C-S-K-P-G-V
L amino acids

LRD peptide 2(1-14)[MIP1 α]: Residues 28-41 of mature hMIP1 α
v-g-p-k-s-c-q-s-s-t-e-f-y-d
D amino acids

LFL peptide 2(1-16)[IL8]: Residues 27-42 of mature hIL8
E-L-R-V-I-E-S-G-P-H-C-A-N-T-E-I
L amino acids

LFL Peptide 2(1-10)[MCP-1]: Residues 28-37 of hMCP-1
S-Y-R-R-I-T-S-S-K-C
L amino acids

LFL peptide 2(10-15)[MCP-1]: Residues 37-42 of hMCP-1
C-P-K-E-A-V
L amino acids

LFL peptide 2(1-5)[MCP-1]: Residues 28-32 of hMCP-1
S-Y-R-R-I
L amino acids

LFL peptide 2(6-10)[MCP-1]: Residues 33-37 of hMCP-1
T-S-S-K-C
L amino acids

LFL peptide 2(1-9)[MIP-1 α]: Residues 28-36 of hMIP-1 α
D-Y-F-E-T-S-S-Q-C
L amino acids

LFL peptide 2(9-14)[MIP-1 α]: Residues 36-41 of hMIP-1 α
C-S-K-P-G-V
L amino acid

LFL Cys₀Ser₁₀Cys₁₆peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1
C-S-Y-R-R-I-T-S-S-K-S-P-K-E-A-V-C
L amino acids

CFL Cys₀Ser₁₀Cys₁₆peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1
C-S-Y-R-R-I-T-S-S-K-S-P-K-E-A-V-C
L amino acids

LRD Cys₀Ser₁₀Cys₁₆peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1
c-v-a-e-k-p-s-k-s-s-t-i-r-r-y-s-c
D amino acids

CRD Cys₀Ser₁₀Cys₁₆peptide 2(1-15)[MCP-1]: Residues 28-42 of hMCP-1
c-v-a-e-k-p-s-k-s-s-t-i-r-r-y-s-c
D amino acids

Sequence	DARC Binding			THP-1 Migration		
	MCP-1	MIP-1 α	SDR-1 α	MCP-1	MIP-1 α	SDR-1 α
SYRRITSSKCPKEAV	350 nM	ns	ns	ns	ns	ns
vaekpckssstirrys	18 μ M	ns	ns	ns	ns	ns
SYRRITSSK	22 μ M	ns	ns	ns	ns	ns
SYRRI	>100 μ M	ns	ns	ns	ns	ns
TSSKC	>100 μ M	ns	ns	ns	ns	ns
CPKEAV	>100 μ M	ns	ns	ns	ns	ns
HLKILMNTPNCAHQIV	19 μ M	10 μ M	40 μ M	7 μ M	7 μ M	7 μ M
DYFETSSQCSPKGV	>100 μ M	ns	ns	ns	ns	ns
vgrkscqssstefyd	>100 μ M	ns	ns	ns	ns	ns
DYFETSSQC	>100 μ M	ns	ns	ns	ns	ns
CSKPGV	>100 μ M	ns	ns	ns	ns	ns

Figure 15

Figure 16

Sequence	Mol Wt.	Duffy Binding ED-50	MCP-1 ED-50	MIP-1 α ED-50	RANTES ED-50	SDF-1 α ED-50	IL-8 ED-50	Other Data
AQDANNAAPVTC	1302	90 μ M	ns	ns	-	ns	ns	
SYRITSSSKCPEKEAV	1725	100nM	ns	ns	-	ns	ns	
vaekpckstirrysv	1725	18 μ M	ns	ns	-	ns	ns	
HEKILATPNCALQIV	1677.3	19 μ M	10 μ M	40 μ M	-	7 μ M	-	
DYFETSSQCSKPGV	1549	>100 μ M	ns	ns	-	ns	ns	
YDPLKSCQSSRTEFVYD	1549	>100 μ M	ns	ns	-	ns	ns	
SYRITSSSKC	1097.4	22 μ M	ns	ns	-	ns	ns	
CPKEAV	645.8	>100 μ M	ns	ns	-	ns	ns	
SYARI	693.9	>100 μ M	ns	ns	-	ns	ns	
TSSKC	525.7	>100 μ M	ns	ns	-	ns	ns	
DYFETSSQCSKPGV	1079.2	>100 μ M	ns	ns	-	ns	ns	
CSKPGV	589.8	>100 μ M	ns	ns	-	ns	ns	

KLK	516.6	>100 μ M					
cq1wkqkpdic	1359	>100 μ M	1nM	-	-	350nM	10nM
cq1wkqkpdic	1448	-	>100 μ M	-	-	-	Note 2
cq1wkqkpdic	1472.2	-	10nM	-	-	-	
SRS	357.3	>100 μ M	>100 μ M	-	-	-	
KKK	609.8	>100 μ M	-	-	-	-	

Note 1: *In Vivo effect- Abolishes macrophages in an *in vivo* rat intradermal study induced by 500ng MCP-1. 300 μ g IV and 10 mg SQ 30 minutes prior to MCP-1.*

Δ Δ Δ (Δ) is associated to Δ Δ Δ (Δ)

Note 2: In same study as Note 1 above, no effect on macrophages seen.

Study Design Table

Group	Animal #	N	RX	RX dose/route T=30 min	Dermal Agonist	Dermal AgonistDo se (ng in 50 ul) T=0	Hour of Sacrifice
1	1,2,3	3	PBS	200 ul: iv 200 ul: SQ-back	PBS LPS MCP-1 MCP-1	0 50 100 500	20-24
2	4,5,6	3	NR58-3.14.3	3 ug: iv 100 ug: SQ back	PBS LPS MCP-1 MCP-1	0 50 100 500	20-24
3	7,8,9	3	NR58-3.14.3	30 ug: iv 1 mg: SQ back	PBS LPS MCP-1 MCP-1	0 50 100 500	20-24
4	10,11,12	3	NR58-3.14.3	300 ug: iv 10 mg: SQ back	PBS LPS MCP-1 MCP-1	0 50 100 500	20-24

Figure 17

United States Patent Application

COMBINED DECLARATION AND POWER OF ATTORNEY

As a below named inventor I hereby declare that: my residence, post office address and citizenship are as stated below next to my name; that

I verily believe I am the original, first and joint inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled: COMPOUNDS AND METHODS TO INHIBIT OR AUGMENT AN INFLAMMATORY RESPONSE.

The specification of which is attached hereto.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, § 1.56 (see page 4 attached hereto).

I hereby claim foreign priority benefits under Title 35, United States Code, § 119/365 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on the basis of which priority is claimed:

No such applications have been filed.

I hereby claim the benefit under 35 U.S.C. § 119(e) of any United States provisional application(s) listed below.

No such applications have been filed.

I hereby claim the benefit under Title 35, United States Code, § 120/365 of any United States and PCT international application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application.

<u>Application Number</u>	<u>Filing Date</u>	<u>Status</u>
08/927,939	September 11, 1997	Pending

I hereby appoint the following attorney(s) and/or patent agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected herewith:

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Drake, Eduardo E.	Reg. No. 40,594	Lacy, Rodney L.	Reg. No. 41,136	Terry, Kathleen R.	Reg. No. 31,884
Dryja, Michael A.	Reg. No. 39,662	Leffert, Thomas W.	Reg. No. 40,697	Viksnins, Ann S.	Reg. No. 37,748
Embreton, Janet E.	Reg. No. 39,665	Lemaire, Charles A.	Reg. No. 36,198	Woessner, Warren D.	Reg. No. 30,440

I hereby authorize them to act and rely on instructions from and communicate directly with the person/assignee/attorney/firm/organization/who/which first sends/sent this case to them and by whom/which I hereby declare that I have consented after full disclosure to be represented unless/until I instruct Schwegman, Lundberg, Woessner & Kluth, P.A. to the contrary.

Please direct all correspondence in this case to Schwegman, Lundberg, Woessner & Kluth, P.A. at the address indicated below:

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Telephone No. (612)373-6900

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like, so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Signature: _____ Date: _____
Suzanne T. Kanaly

Full Name of inventor:

Citizenship: _____ Residence: _____
Post Office Address: _____

Signature: _____ Date: _____

§ 1.56 Duty to disclose information material to patentability.

(a) A patent by its very nature is affected with a public interest. The public interest is best served, and the most effective patent examination occurs when, at the time an application is being examined, the Office is aware of and evaluates the teachings of all information material to patentability. Each individual associated with the filing and prosecution of a patent application has a duty of candor and good faith in dealing with the Office, which includes a duty to disclose to the Office all information known to that individual to be material to patentability as defined in this section. The duty to disclose information exists with respect to each pending claim until the claim is canceled or withdrawn from consideration, or the application becomes abandoned. Information material to the patentability of a claim that is canceled or withdrawn from consideration need not be submitted if the information is not material to the patentability of any claim remaining under consideration in the application. There is no duty to submit information which is not material to the patentability of any existing claim. The duty to disclose all information known to be material to patentability is deemed to be satisfied if all information known to be material to patentability of any claim issued in a patent was cited by the Office or submitted to the Office in the manner prescribed by §§ 1.97(b)-(d) and 1.98. However, no patent will be granted on an application in connection with which fraud on the Office was practiced or attempted or the duty of disclosure was violated through bad faith or intentional misconduct. The Office encourages applicants to carefully examine:

- (1) prior art cited in search reports of a foreign patent office in a counterpart application, and
- (2) the closest information over which individuals associated with the filing or prosecution of a patent application believe any pending claim patentably defines, to make sure that any material information contained therein is disclosed to the Office.

(b) Under this section, information is material to patentability when it is not cumulative to information already of record or being made of record in the application, and

- (1) It establishes, by itself or in combination with other information, a prima facie case of unpatentability of a claim; or
- (2) It refutes, or is inconsistent with, a position the applicant takes in:
 - (i) Opposing an argument of unpatentability relied on by the Office, or
 - (ii) Asserting an argument of patentability.

A prima facie case of unpatentability is established when the information compels a conclusion that a claim is unpatentable under the preponderance of evidence, burden-of-proof standard, giving each term in the claim its broadest reasonable construction consistent with the specification, and before any consideration is given to evidence which may be submitted in an attempt to establish a contrary conclusion of patentability.

(c) Individuals associated with the filing or prosecution of a patent application within the meaning of this section are:

- (1) Each inventor named in the application;
- (2) Each attorney or agent who prepares or prosecutes the application; and
- (3) Every other person who is substantively involved in the preparation or prosecution of the application and who is associated with the inventor, with the assignee or with anyone to whom there is an obligation to assign the application.

(d) Individuals other than the attorney, agent or inventor may comply with this section by disclosing information to the attorney, agent, or inventor.